

UNIVERSITY OF WOLLONGONG

DOCTORAL THESIS

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**Constraining natural contributions to  
tropospheric ozone production over  
Australia and selected areas of the  
Southern Ocean**

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*Author:*  
Jesse GREENSLADE

*Supervisors:*  
Jenny Fisher  
Clare Murphy

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in the*

Centre for Atmospheric Chemistry  
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December 13, 2019



I, Jesse GREENSLADE, declare that this thesis titled, "Constraining natural contributions to tropospheric ozone production over Australia and selected areas of the Southern Ocean" and the work presented in it are my own. I confirm that:

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- Where I have consulted the published work of others, this is always clearly attributed.
- Where I have quoted from the work of others, the source is always given. With the exception of such quotations, this thesis is entirely my own work.
- I have acknowledged all main sources of help.
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Date: 1/8/2019



# Abstract

Ozone in the troposphere is a toxic pollutant that causes respiratory and agricultural damage. The two main sources of tropospheric ozone are chemical production, and transport from the stratosphere. Australia lacks the extensive in-situ measurement sites (or network) required to fully attribute ozone concentrations to these different sources within either regional or urban areas.<sup>mh</sup> The primary source is chemical production, which Chemical production<sup>mh</sup> can occur following biogenic emissions of volatile organic compounds (VOCs) when they mix with polluted urban air. Most tropospheric ozone is formed through chemical reactions involving nitrogen oxides, the hydroxyl radical, and VOCs. Most emitted VOCs are of biogenic origin, and the primary biogenic VOC emitted to the atmosphere from land is isoprene; however, estimates of isoprene emission rates are highly uncertain. The second most important source of tropospheric ozone is the stratosphere, which occasionally mixes into the troposphere bringing ozone-rich air masses down towards the Earth's surface.<sup>mh</sup> Transport of ozone from the stratosphere is also<sup>mh</sup> uncertain, and difficult to measure. These uncertainties affect atmospheric chemistry models, reducing confidence in modelled atmospheric processes such as radiative forcing and air quality forecasting.

This thesis has three aims: to quantify formaldehyde amounts over Australia observed by satellite using a global chemical transport model (GEOS-Chem), to determine Australian isoprene emissions using modelled isoprene-to-formaldehyde yields along with satellite formaldehyde amounts, and to attribute ozone in the troposphere to the contributions from chemical production and stratospheric transport. Model output and observations are combined for each aim in this thesis.

Formaldehyde measurement records from satellite instruments can be used to estimate isoprene emission rates. Satellite formaldehyde measurements require modelled a priori vertical profiles of concentration. Corrections are required to remove the influence of this a priori profile when comparing satellite products against models or other measurements. In this thesis, formaldehyde measurements were recalculated in two different ways (removing the effects from the old a priori), with both recalculation techniques and outcomes analysed. The method used to recalculate vertical column amounts significantly affected the outcome. Measurements affected by anthropogenic and pyrogenic sources were filtered out using other satellite products, and then satellite pixels were gridded into a daily dataset. These data were formed to be used with modelled isoprene-to-formaldehyde yields in order to estimate isoprene emissions.

Isoprene is predominantly emitted by trees and shrubs, and Eucalypts are potentially very high emitters. Isoprene oxidises in the atmosphere to form formaldehyde, which has a sufficiently short lifetime in the atmosphere to establish chemical equilibrium. In this thesis, aA<sup>mh</sup> linear model was created using modelled isoprene emissions and associated formaldehyde enhancements over Australia. This model was applied

to satellite formaldehyde to create a new estimate of Australian isoprene emissions. Isoprene emissions from Australian forests were found to be substantially lower than previous estimates have predicted. **Summer (peak season) reductions averaged at ~ 50% and varied spatially.**<sup>mh</sup> This leads to an overall lowering of surface background ozone concentrations of approximately 5%.

In order to examine potential stratospheric sources of tropospheric ozone, ~~The second most important source of tropospheric ozone is the stratosphere, which occasionally mixes into the troposphere bringing ozone-rich air masses down towards the Earth's surface. In this thesis,~~<sup>mh</sup> an analysis of the local weather patterns and ozone seasonality was performed, showing most stratosphere to troposphere transport occurs during low pressure frontal weather systems. This work provides a novel technique using a mathematical (Fourier band-pass) filter on ozonesonde profiles for estimation and quantification of tropospheric ozone transported from the stratosphere. An estimate encompassing three measurement stations over the Southern Ocean near Australia of about  $7.2 \times 10^{17}$  molec cm<sup>-2</sup> per year was derived.

Overall, this thesis improves knowledge of two important natural sources of Australian tropospheric ozone concentrations by examining and estimating isoprene emissions and by characterising stratospheric intrusions based on ozonesondes. **Background ozone concentrations are the basis for ozone pollution and air quality metrics. How these concentrations are affected by over-represented isoprene emissions and by transport from the stratosphere is now better understood for Australia and the nearby Southern Ocean.**<sup>mh</sup>

## *Acknowledgements*

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# List of Abbreviations

<b>ACCMIP</b>	Atmospheric Chemistry and Climate Model Inter-comparison Project
<b>AAOD</b>	Aerosol Absorption Optical Depth
<b>AMF</b>	Air Mass Factor
<b>AOD</b>	Aerosol Optical Depth
<b>BVOC</b>	Biogenic Volatile Organic Compound
<b>CCD</b>	Charged Coupled Device spectrometer
<b>CPC</b>	Climate Prediction Center
<b>CTM</b>	Chemical Transport Model
<b>DOAS</b>	Differential Optical Absorption Spectroscopy
<b>ECMWF</b>	European Centre for Medium-range Weather Forecasts
<b>EDGAR</b>	Emission Database for Global Atmospheric Research
<b>ERA-I</b>	ECMWF ReAnalysis (Interim)
<b>GC-FID</b>	Gas Chromatographer Flame Ionisation Detector
<b>FTIR</b>	Fourier transform Infra-Red spectrometer
<b>GEOS</b>	Goddard Earth Observing System
<b>GMAO</b>	Global Modeling and Assimilation Office
<b>GOME</b>	Global Ozone Monitoring Experiment
<b>GPH</b>	GeoPotential Height
<b>HEMCO</b>	Harvard-NASA Emissions Component
<b>IQR</b>	Inter-Quartile Range
<b>LAI</b>	Leaf Area Index
<b>LT</b>	Local Time
<b>OMHCHO</b>	OMI satellite HCHO product
<b>OMHCHORP</b>	OMI satellite HCHO product re-processed
<b>OMI</b>	Ozone Monitoring Instrument
<b>MEGAN</b>	Model of Emissions of Gases and Aerosols from Nature
<b>MUMBA</b>	Measurements of Urban, Marine, and Biogenic Air
<b>NCO</b>	National Computing Infrastructure
<b>NDACC</b>	Network for the Detection of Atmospheric Composition Change
<b>NH</b>	Northern Hemisphere
<b>NMVOC</b>	Non-Methane Volatile Organic Compound
<b>(S,P)OA</b>	(Secondary, Primary) Organic Aerosols
<b>OMR</b>	Ozone Mixing Ratio
<b>PAN</b>	PeroxyAcetyl Nitrate
<b>PFT</b>	Plant Functional Type
<b>PM</b>	Particulate Matter
<b>PTR-MS</b>	Proton-Transfer-Reaction Mass spectrometer
<b>PV</b>	Potential Vorticity
<b>RA</b>	Row Anomaly

<b>RF</b>	Radiative Forcing
<b>RMA</b>	Reduced Major Axis
<b>RMSA</b>	Root Mean Square Error
<b>RSC</b>	Reference Sector Correction
<b>RTM</b>	Radiative Transfer Model
<b>SAO</b>	Smithsonian Astrophysical Observatory
<b>SH</b>	Southern Hemisphere
<b>SHADOZ</b>	Southern Hemisphere Additional OZonesonde
<b>SPS(1,2)</b>	Sydney Particulate Studies
<b>STT</b>	Stratosphere to Troposphere Transport
<b>SZA</b>	Solar Zenith Angle
<b>TOA</b>	Top Of the Atmosphere
<b>TOMS</b>	Total Ozone Mapping Spectrometer
<b>VOC</b>	Volatile Organic Compounds
<b>UCX</b>	Universal tropospheric-stratospheric Chemistry eXtension
<b>UV-Vis</b>	Ultraviolet and visible
<b>VCC</b>	Vertical Column Corrected
<b>VMR</b>	Vertical Mixing Ratio
<b>WOUDC</b>	World Ozone and Ultraviolet Data Centre

## Chapter 1

# Introduction and Literature Review

### 1.1 The atmosphere

The atmosphere is made up of gases held to the earth's surface by gravity. These gases undergo transport on all scales, from barbecue smoke being blown about the garden, to smoke plumes from forest fires travelling across the world and depositing in the Antarctic snow. They take part in innumerable chemical reactions along the way, largely driven by solar input and interactions with each other. Many gases are emitted into the atmosphere by soil, trees, factories, cars, seas and oceans. They are also deposited back to the surface both directly and in rainfall.

The atmosphere is made up of nitrogen ( $N_2$ :  $\sim 78\%$ ), oxygen ( $O_2$ :  $\sim 21\%$ ), and argon ( $Ar$ :  $\sim 1\%$ ), along with water ( $H_2O$ ) and *trace gases* (those that make up less than 1% of the atmosphere). Atmospheric  $H_2O$  content can be as high as 4% depending on local conditions. Beyond these major constituents the atmosphere has a vast number of trace gases, including carbon dioxide ( $CO_2$ :  $\sim 0.4\%$ ), ozone ( $O_3$ :  $.000001\%$  to  $0.001\%$ ), and methane ( $CH_4$ :  $\sim 0.00018\%$ ) (*Earth System Research Laboratory; Global Monitoring Division*; Brasseur and Jacob 2017, Ch. 2). Trace gases in the atmosphere can have a large impact on conditions for life on earth. They combine, break apart, and react with each other affecting all surface ecosystems upon which life depends.

One important trace gas is ozone, as it directly affects the climate, human health, and ecosystem productivity. The ozone budget (production, loss, and transport) is relatively uncertain over Australia. This thesis focuses on ozone in the troposphere over Australia and sites in the nearby Southern Ocean. It also estimates emissions of isoprene, one of the important precursors to tropospheric ozone production. This chapter provides background on the structure and composition of the atmosphere and introduces the key atmospheric species examined in this thesis, as well as relevant techniques used to measure and model chemistry in the atmosphere.

#### 1.1.1 Structure

Most of the atmosphere ( $\sim 85\%$ ) is within 10 km of the earth's surface. This is due to gravity, which causes air pressure to decrease logarithmically with altitude. Any entity is subjected to the weight of the air above it, and the structure of the atmosphere is driven by this pressure.

The atmosphere extends above the earth's surface to the edges of space. This is split into various layers, defined by the *lapse rate*: the decrease in temperature ( $T$ ) with

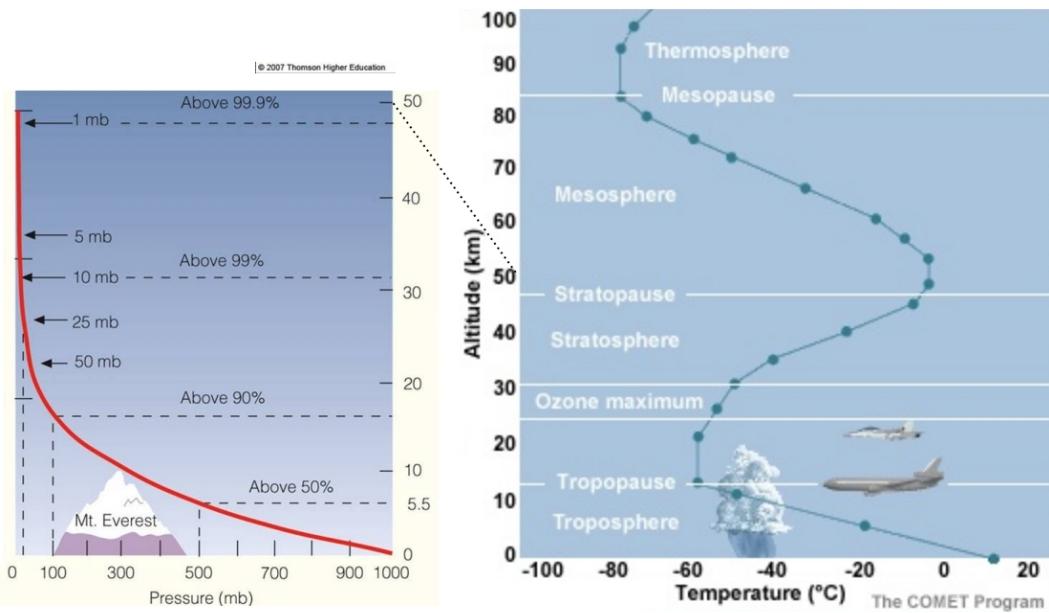


FIGURE 1.1: Pressure (red) logarithmically decreasing, shown with percentage of atmosphere below at several points. Temperature (green) changes throughout the atmosphere. Figure modified from <https://climate.ncsu.edu/edu/Structure>.

increasing altitude ( $z$ ), or  $\frac{-dT}{dz}$ . Figure 1.1 shows the pressure and temperature profiles against altitude through the atmosphere. The first layer is the troposphere, which extends to roughly 10 km and is characterised by positive lapse rate (or decreasing temperature with altitude). At the top of the troposphere (the tropopause) the temperature stops decreasing, and then the stratosphere is defined by a negative lapse rate. This is due to ultraviolet radiation being absorbed by ozone, and leads to a very vertically stable environment.

In addition to these atmospheric layers, the troposphere can be subset into the *boundary layer* and the *free troposphere*. The *boundary layer* is the lowest layer and involves increased atmospheric mixing due to ground heating and friction effects. It generally extends from the surface up to 200 m - 1000 m, above which the ground effects have fewer direct impacts. The *free troposphere* is the remainder of the troposphere, where trace gas concentrations are more affected by transport and chemistry. Transported trace gases (and particulates) can come from the stratosphere or local to global scale winds and jet streams.

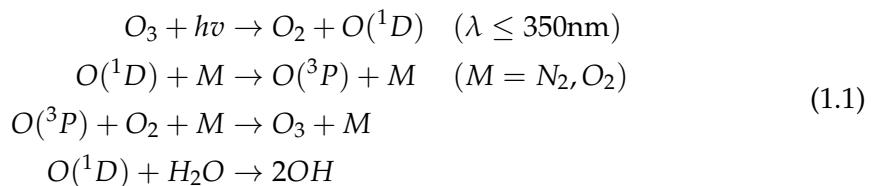
### 1.1.2 Composition and chemistry

There are myriad trace gases in the atmosphere, emitted by plants, animals, earth, and water. These gases react with one another and over time they either deposit back onto the earth or form more stable compounds such as carbon dioxide ( $\text{CO}_2$ ). Oxidation

and photolysis (the process of being broken apart by photons) are the two main processes whereby compounds are broken down in the atmosphere. Products formed in these reactions are sometimes called child products.

Hydroxyl ( $\text{OH}$ ) and hydroperoxyl ( $\text{HO}_2$ ) radical<sup>b</sup> concentrations largely determine the oxidative capacity of the atmosphere. The OH radical drives many processes in the atmosphere, especially during the day when it is produced by the photolysis of ozone (Atkinson 2000). OH is a key species that reacts with nearly all the organic compounds in the troposphere, with only a few exceptions (Atkinson 2000). Over land, isoprene ( $\text{C}_5\text{H}_8$ ) and monoterpenes ( $\text{C}_{10}\text{H}_{16}$ ) account for 50% and 30% of the OH reactivity respectively (Fuentes et al. 2000).

Since radicals are involved in all oxidative chemistry in the atmosphere it is important for models to accurately represent them. This is difficult as they are coupled with so many other species and measurements of OH are not readily available on a global scale. OH formation in the clean troposphere is driven by ozone photolysis, as excited oxygen atoms ( $\text{O}({}^1\text{D})$ ) are created and react with water to form OH, shown in reaction sequence 1.1 (Atkinson 2000; Atkinson and Arey 2003):



where  $h\nu$  represents radiation and M is an inert molecule. This shows that some of the  $\text{O}({}^1\text{D})$  recycles back to ozone, while some forms OH. In the late 1990s it was thought that OH radicals were formed exclusively from photolysis of  $\text{O}_3$ , HONO, HCHO, and other carbonyls ( $\text{R}_2\text{C=O}$ ) (Atkinson 2000). It has been shown since that OH is recycled in various processes. For example isoprene ( $\text{C}_5\text{H}_8$ ) was thought to be a sink of OH until it was shown by Paulot et al. (2009b) that the radicals are recycled. This recycling process is discussed in more detail in section 1.3.4.

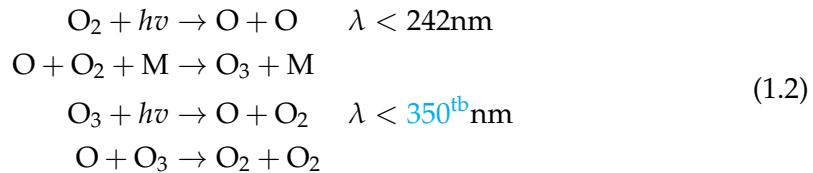
## 1.2 Ozone

Ozone ( $\text{O}_3$ ) is an important greenhouse gas and oxidant. It is mostly located in the stratosphere and prevents much of the shorter wavelength solar radiation from reaching the earth's surface. Ozone in the troposphere is less beneficial, leading to health issues, radiative forcing, and crop death (Stevenson et al. 2013). Understanding and accurately portraying ozone concentrations in the troposphere is important to allow accurate predictions of future climate. This will become even more important as climate change alters the atmosphere (Hegglin and Shepherd 2009).

### 1.2.1 Stratospheric ozone

In the stratosphere, ozone production is driven by the Chapman mechanism, as high energy radiation (with wavelengths  $\lambda < 242$  nm) photolyses the molecular oxygen ( $\text{O}_2$ ) in the atmosphere (Brasseur and Jacob 2017, Chapter 3, section 2). The Chapman

mechanism involves several reactions that lead to rough equilibrium of O, O<sub>2</sub>, O<sub>3</sub> and pressure, as follows:



The high energy photons ( $\lambda < 242$  nm) are present from the top of the atmosphere but are mostly removed before reaching the troposphere as their energy is used to split the O<sub>2</sub> molecules. The lifetime of O against loss by O<sub>2</sub> is less than a second in the troposphere, and produced O<sub>3</sub> quickly returns to O and O<sub>2</sub>, as *lower energy ( $\lambda < 350$  nm) photons are abundant*~~low energy ( $\lambda < 1180$  nm) photons and M are abundant~~<sup>tb</sup>. The reduced light penetration towards the surface, in addition to the logarithmic increase in atmospheric pressure (which affects M abundance) *results in a vertical profile of ozone that peaks in the stratosphere*~~.drives the vertical profile of ozone into what is called the ozone layer.~~<sup>tb</sup> This is a layer of relative ozone abundance within the stratosphere *called the ozone layer*<sup>tb</sup>.

Satellite based measurements of ozone were instrumental in identifying a gap in the ozone layer (the ozone hole), where ozone concentrations were unexpectedly low. Since the Montreal Protocol on Substances that Deplete the Ozone Layer was established in August 1987, and ratified in August 1989, new satellites and measurement stations were set up to monitor ozone in the stratosphere. Detecting ozone from the surface up to the top of the stratosphere requires techniques such as remote sensing and ozonesonde releases. Ozonesondes are weather balloons (with attached ozone detectors) that detect ozone concentrations up to the mid stratosphere (~ 30 km), providing a vertical profile over a single location. Ozonesondes have been released periodically for decades over some cities, allowing long term ozone concentration profile analysis (e.g., Brinksma et al. 2002). A small network of ozonesonde release sites (including Davis, Macquarie Island, and Melbourne) is available from the world ozone and ultraviolet radiation data centre <http://woudc.org/data/explore.php> and is used in Chapter 4 to examine stratospheric impacts on tropospheric ozone (see Section 2.2.3.2 for more info on these ozonesondes).

## 1.2.2 Tropospheric ozone

Ozone in the lower atmosphere is a serious hazard that causes health problems (Hsieh and Liao 2013), causes billions of dollars of damage to agricultural crops (Avnery et al. 2013; Yue et al. 2017), and increases the rate of climate warming (Myhre and Shindell 2013). Around 5 to 20 percent of all air pollution related deaths are due to ozone (Monks et al. 2015), which translates to roughly 800 thousand deaths per year (Lelieveld et al. 2013). In the short term, ozone concentrations of ~50-60 ppbv over eight hours or ~80 ppbv over one hour are agreed to constitute a human health hazard (Ayers and Simpson 2006; Lelieveld et al. 2009). Long term exposure causes problems with crop loss and ecosystem damage (Ashmore, Emberson, and Murray Frank 2003), and concentrations may get worse in the future (Lelieveld et al. 2009; Stevenson et al.

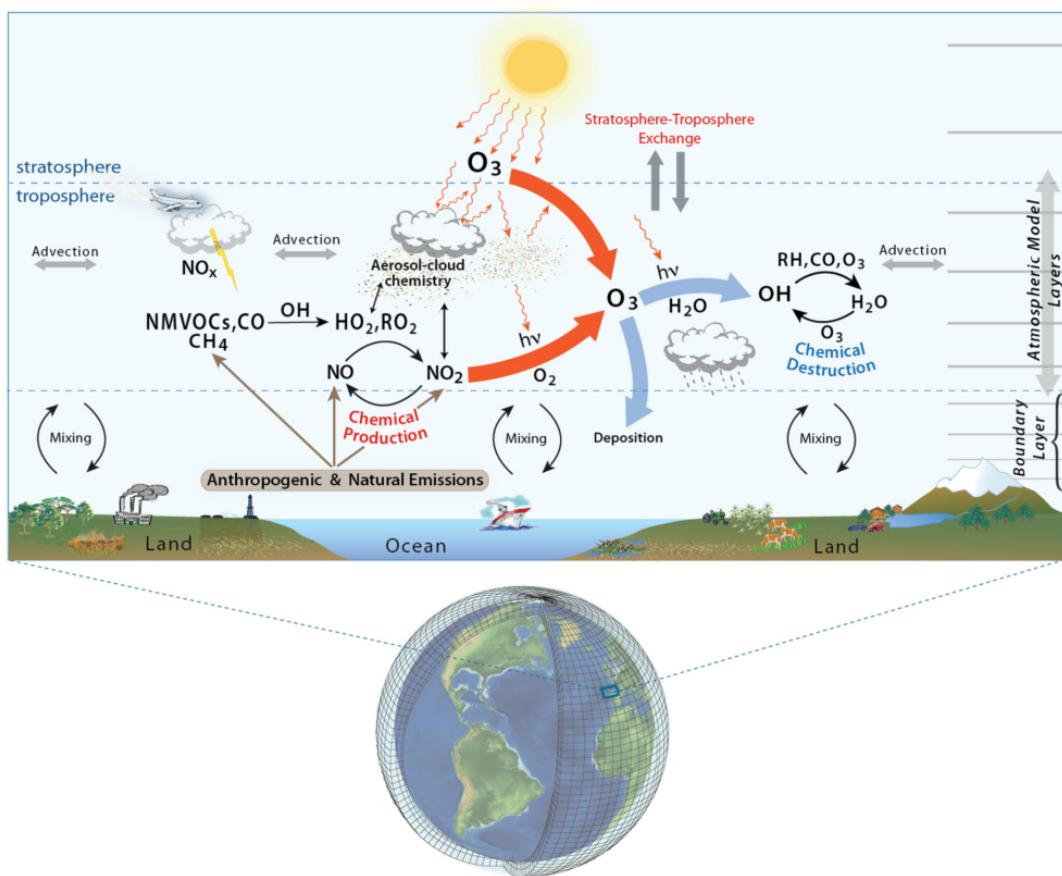


FIGURE 1.2: Tropospheric ozone processes, Figure 1 in Young et al. (2017). DOI: <https://doi.org/10.1525/elementa.265.f1>.

2013). For example, future tropospheric ozone enhancements are projected to drive reductions in global crop yields equivalent to losses of up to \$USD<sub>2000</sub> 35 billion (US dollars in the year 2000) per year by 2030 (Avnery et al. 2013), along with detrimental health outcomes equivalent to ~\$USD<sub>2000</sub> 11.8 billion per year by 2050 (Selin et al. 2009). Recently Yue et al. (2017) showed that the net effect of near-surface ozone is an approximately 14% decrease in net primary productivity in China. They also state that in order to wind back most of this productivity decrease, drastic measures are required.

Figure 1.2, reproduced from Young et al. (2017), shows a summary of the major processes and emissions affecting tropospheric ozone. This thesis focuses on improving the highly uncertain natural emissions of non-methane volatile organic compounds (NMVOCs) from Australia, and estimating how much ozone is transported down from the stratosphere.

Generally there are two main drivers of tropospheric ozone concentrations: transport from the stratosphere and chemical production due to emissions of precursors. Tropospheric ozone is lost via chemical destruction and dry deposition, estimated to be  $4700 \pm 700 \text{ Tg yr}^{-1}$  and  $1000 \pm 200 \text{ Tg yr}^{-1}$ , respectively (Stevenson et al. 2006; Young et al. 2017). The main loss channel is through photolysis and collisions, and

leads to OH production (equation 1.1).

### 1.2.2.1 Stratosphere to troposphere transport

Historically, ozone transported down from the stratosphere was thought to contribute 10-40 ppb to tropospheric ozone levels, making up 50% of tropospheric concentrations (Atkinson 2000; Stohl et al. 2003). The proportion was revised down to around 10% over the years as measurement and modelling campaigns improved our<sup>mh</sup> understanding of global scale transport, mixing, and chemistry (Guenther et al. 2006; Monks et al. 2015). Intrusions of stratospheric air into the troposphere are often called Stratosphere to Troposphere Transport (STT) events. Although most tropospheric ozone comes from production, STT enhancements of ozone are measurable and can be regionally important (e.g., Jacobson and Hansson 2000; Lelieveld et al. 2009; Kuang et al. 2017). Additionally, upper tropospheric ozone can be transported long distances (Cooper et al. 2004), affecting measurements far downwind of where stratospheric mixing may be taking place. An analysis of the Atmospheric Chemistry and Climate Model Inter-comparison Project simulations showed STT enhances tropospheric ozone by  $540 \pm 140 \text{ Tg yr}^{-1}$  (Young et al. 2013), equivalent to  $\sim 11\%$  of the tropospheric ozone column (Monks et al. 2015).

Ozone transported to the troposphere from the stratosphere can occur through diffusion (relatively slowly) or direct mixing (as STT). STT often occur as tongues of stratospheric air that descend into the troposphere before becoming detached. These can be due to low pressure systems and jet streams (Sprenger, Croci Maspoli, and Wernli 2003; Alexander et al. 2013). It is possible to model this process and estimate how much ozone is being transported by STT (e.g., Young et al. 2013; Ojha et al. 2016). Model based estimates require validation against actual measurements, such as those from ozonesondes or satellites. It has been estimated that climate change will lead to increased STT through acceleration of the Brewer Dobson circulation (Hegglin and Shepherd 2009), a large scale transport system that affects the structure and composition of the atmosphere, and meteorology, in the tropics. Hegglin and Shepherd (2009) estimated that ozone transport would increase by 23% globally by 2095 (relative to 1965),  $\sim 30 \text{ Tg yr}^{-1}$  in the southern hemisphere and  $\sim 121 \text{ Tg yr}^{-1}$  in the northern hemisphere.

STT mostly impacts the upper troposphere, although some areas are impacted down to the surface. Over both ocean and land STT can lead to seasonal enhancements of upper and lower tropospheric ozone concentrations (Lin et al. 2015; Liu et al. 2017a; Kuang et al. 2017). A confounding factor is that stratospheric based ozone enhancement can be transported horizontally, and is not confined to the area where intrusion into the troposphere occurred. Understanding of STT needs to be improved to allow source attribution of the causes of local ozone enhancements (Lin et al. 2015).

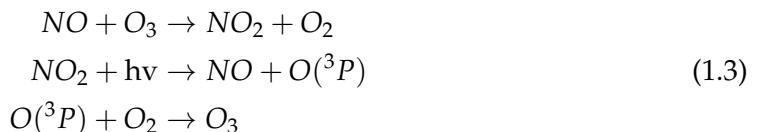
### 1.2.2.2 Chemical production

Tropospheric ozone concentrations are enhanced by ozone precursor emissions; including NO, NO<sub>2</sub>, CO, and VOCs such as isoprene (Atkinson 2000; Young et al. 2013; Marvin et al. 2017). Ozone predictions are uncertain and changing climate affects

transport, deposition, destruction, and natural precursor emissions. All of these processes are tightly coupled and difficult to accurately model, as they depend on uncertain assumptions such as CO<sub>2</sub> dependency (Young et al. 2013). Despite significant advances over the prior decades, there remain large uncertainties about ozone precursors in the troposphere (Mazzuca et al. 2016). Tropospheric ozone is regulated by NO and NO<sub>2</sub> concentrations, which form an equilibrium (Cape 2008; Young et al. 2017). At all scales, pyrogenic (fire) and anthropogenic (man-made) emissions can be important; however, biogenic VOC emissions are a nearly constant source of potential production during the day. Smoke plumes from biomass burning can carry ozone precursors, creating higher ozone concentrations downwind of the plume's source. Emissions of precursors from large cities (primarily from traffic and power production) can impact ozone concentrations. These impacts are not always straightforward due to the nonlinear relationship between ozone and its precursors. Recently, modelled ozone concentrations have been shown to be most sensitive to NO<sub>x</sub> ( $\equiv$  NO<sub>2</sub> + NO) sources (such as lightning and car exhaust emissions) and isoprene emissions (Christian et al. 2018).

NO<sub>x</sub> is an important chemical family in the atmosphere, which interacts with ozone and regulates the atmospheric oxidative capacity. NO<sub>x</sub> and VOC emissions affect the tropospheric ozone equilibrium and can lead to enhanced ozone formation, shown in figure 1.2. NO<sub>x</sub> compounds are short-lived, with emissions from power generation and transport being the main driver of concentrations (Delmas, Serca, and Jambert 1997). NO<sub>x</sub> is removed primarily by conversion to nitric acid (HNO<sub>3</sub>) or alkyl nitrates (RONO<sub>2</sub>) followed by wet or dry deposition (Ayers and Simpson 2006; Romer Present, Zare, and Cohen 2019). Ozone in rural areas is often higher than in populous cities, due to titration (removal) of ozone by NO in areas with high anthropogenic emissions (Cooper, Gilge, and Shindell 2014; Monks et al. 2015).

As discussed above, STTs are responsible for  $\sim 11\%$  of the tropospheric column of ozone, with the remainder produced photochemically. A recent summary by Young et al. (2017) estimated ozone production and loss in the troposphere to be  $\sim 4900 \text{ Tg yr}^{-1}$ , and  $\sim 4500 \text{ Tg yr}^{-1}$  respectively. These numbers are at the global scale, and it should be noted that meteorology and topography can play dominant roles at local to regional spatial scales (e.g., Kuang et al. 2017). The main processes involved are shown in figure 1.2, with ozone and NO<sub>x</sub> concentrations regulated by the following reactions (Sillman 1999; Atkinson 2000):



VOC emissions can alter this process, essentially replacing ozone in the first stage of the above equation. The process without and with the influence of VOCs (panel A and B respectively) is summarised in Figure 1.3. Depending on local atmospheric conditions, ozone production can be limited by the presence of VOC or NO<sub>x</sub> concentrations, and this is often referred to as the VOC limited or NO<sub>x</sub> limited regime.

Net formation or loss of O<sub>3</sub> is determined by interactions between VOCs, NO<sub>x</sub>, and HO<sub>x</sub>, and is a complicated system of positive and negative feedbacks (Atkinson 2000).

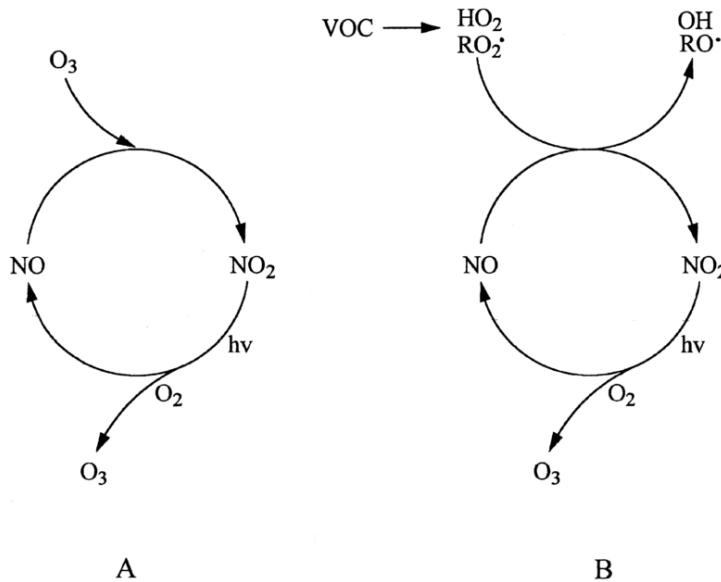


FIGURE 1.3: NO - NO<sub>2</sub> - O<sub>3</sub> photoequilibrium cycle without and with (A, B respectively) influence from VOCs. Figure reproduced from Atkinson (2000).

Figure 1.4 shows an example of this non-linear relationship between NO<sub>x</sub>, VOCs, and ozone production as modelled in Mazzuca et al. (2016). Essentially increasing NO<sub>x</sub> and VOC concentrations will increase ozone production; however, increasing one or the other may or may not increase production depending on current conditions. This is a short time scale relationship and ozone production can be more or less sensitive to VOCs at different hours of the day (Mazzuca et al. 2016). It is important to correctly determine the precursor concentrations in order to estimate ozone levels and production. This non-linear relationship is examined in more detail in the following section (1.3).

### 1.3 Volatile Organic Compounds

The least well understood precursors to tropospheric ozone production belong to a subclass of organic compounds. Organic compounds are a class of chemicals whose molecules contain carbon, with the exception of a few compounds such as carbides, carbonates, and simple oxides of carbon and cyanide. Organic compounds can be categorised based on their vapour pressure, which is the tendency of a liquid or solid to vaporise. Compounds with high vapour pressures at standard temperature are classed as volatile (volatile organic compounds: VOC), and are likely to vaporise at surface pressure. Atmospheric organic compounds are legion and differ by orders of magnitude with respect to their fundamental properties, such as volatility, reactivity, and cloud droplet formation propensity. **These properties are largely dictated by the**

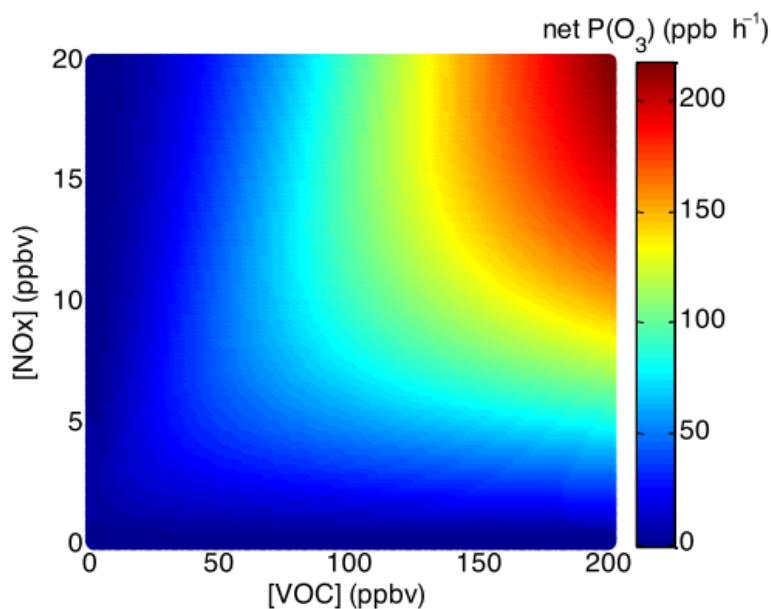


FIGURE 1.4: Dependence of ozone production rate on  $\text{NO}_x$  and VOC concentrations reproduced from Mazzuca et al. (2016).

chemical makeup and structure of the individual compounds. A compound's atmospheric lifetime is strongly related to its reactivity (and the concentration of reactants), with more reactive compounds having shorter atmospheric lifetimes.<sup>tb</sup>

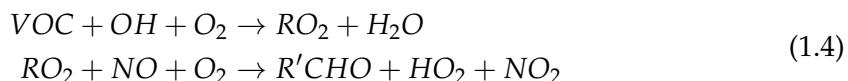
VOCs have vapour pressure greater than  $10^{-5}$  atm, and are mostly generated naturally by plants, which emit around  $1000 \text{ Tg yr}^{-1}$  (Guenther et al. 1995; Glasius and Goldstein 2016). Due to their high volatility these compounds generally exist in the gas phase. Organic compounds with a lower volatility are classed as semi-volatile (SVOCs: vapour pressure between  $10^{-5}$  and  $10^{-11}$  atm) and are found in both gas and particle phase depending on temperature and pressure. Organic compounds with even lower vapour pressure are generally found in the particle phase (Glasius and Goldstein 2016). Plants contain tens of thousands of organic compounds, with fewer than 40 having high enough volatility to be emitted (Guenther et al. 2000). Gas phase compounds with higher vapour pressures can be oxidised into lower vapour pressure products that partition between gas and particle phase. For instance VOCs are oxidised in the atmosphere (mainly by OH), forming HCHO, O<sub>3</sub>, CO<sub>2</sub> and many other species.

Globally, biogenic sources dominate emissions of VOCs (Guenther et al. 1995; Kanakidou et al. 2005; Millet et al. 2008; Guenther et al. 2012), although anthropogenic and pyrogenic sources can also be important in specific environments (e.g., Zhu et al. 2014a; Kwon et al. 2017).<sup>tb</sup> Understanding the drivers of trends in biogenic VOC emissions (BVOCs) is required in order to estimate future carbon fluxes, changes in the water cycle, ozone production, air quality, and other climate responses (Yue, Unger, and Zheng 2013). For example, in China over the last 20 years, anthropogenic emissions of VOCs have been increasing while biogenic VOC emissions have decreased, due to rapid economic

growth and lower annual temperatures (Stavrakou et al. 2014; Kwon et al. 2017). Understanding the drivers of trends in biogenic VOC emissions (BVOCs) is required in order to estimate future carbon fluxes, changes in the water cycle, ozone production, air quality, and other climate responses (Yue, Unger, and Zheng 2015).<sup>tb</sup>

Methane ( $\text{CH}_4$ ) is one of the more abundant VOCs, however it is often classified separately from non-methane VOCs (NMVOCs). NMVOCs include alkanes, alkenes, and aromatic hydrocarbons, with isoprene (an alkene) being the most abundant (Guenther et al. 1995). Methane is relatively long lived (years) and is well mixed in the atmosphere, while other VOCs have shorter lifetimes and therefore their concentrations are more spatially variable. VOCs with short lifetimes are generally only found in high concentrations near their emission sources.

In areas with high VOC concentrations, ozone production may be enhanced through the following reaction sequence (Sillman 1999):



with R and R' representing organic species. The reactions of VOCs with OH convert NO to  $\text{NO}_2$ , which leads to greater ozone formation as  $\text{NO}_2$  production is bypassed in reaction 1 of Equation 1.3.

One aspect associated with VOC emissions is the production of aerosols. Aerosols are suspended particulates and liquid compounds in the atmosphere, often called particulate matter (PM). PM in the atmosphere is a major problem, causing an estimated 2-3 million deaths annually (Avnery et al. 2013; Hoek et al. 2013; Krewski et al. 2009; Silva et al. 2013; Lelieveld et al. 2015). Fine particulate matter ( $\text{PM}_{2.5}$ ) penetrates deep into the lungs and is detrimental to human health. Some PM comes from small organic aerosols (OA) that are directly emitted to the atmosphere in the particulate phase, referred to as primary OA (POA).

A substantial amount of PM is due to gaseous organic compounds transforming into what is known as secondary OA (SOA) (Atkinson 2000; Kanakidou et al. 2005; Kroll and Seinfeld 2008). Formation of SOA is generally due to VOC oxidation and subsequent reactions, while removal from the atmosphere is largely due to wet or dry deposition, and cloud scavenging (Kanakidou et al. 2005). Most of the tropospheric SOA comes from biogenic precursors, and the evidence for this has grown over the last two decades (Guenther et al. 1995; Kanakidou et al. 2005; Guenther et al. 2012). Improved concentration estimates of these precursors requires a better understanding of their emissions, which is one of the foci in this thesis.

VOCs are removed mainly by photolysis and oxidation by OH, forming alkyl radicals ( $\text{R}'$ ). Additional losses are caused by wet and dry deposition, reaction with  $\text{NO}_3$ , and ozonolysis (at night time or in polluted areas) (Atkinson and Arey 2003; Brown et al. 2009). Deposition only accounts for a small fraction of the VOC loss (Atkinson and Arey 2003).

### 1.3.1 Emissions

VOC emissions are classified as either biogenic (BVOC), anthropogenic, or pyrogenic. Global NMVOC emissions are estimated to come from (approximately) 85% biogenic,

13% anthropogenic, and 3% pyrogenic sources respectively (Kanakidou et al. 2005; Kefauver, Filella, and Peñuelas 2014). This thesis mostly focuses on biogenic emissions, with influences from pyrogenic and anthropogenic emissions removed (Section 3.2.3) and not discussed further.<sup>tb</sup> The ocean also plays a role in VOC emissions, with the Oceanic Niño Index showing positive VOC emission anomalies associated with neighbouring countries (Stavrakou et al. 2014). Due to the lack of in situ ground based measurements, estimates of VOC emissions are uncertain, with large scale extrapolation required (Millet et al. 2006).

There is ten times as much mass of NMVOCs emitted from natural sources as from anthropogenic sources (Guenther et al. 2006; Kanakidou et al. 2005; Millet et al. 2006). Major emitters are broadleafs (notably Eucalyptus), and shrubs (Guenther et al. 2006; Arneth et al. 2008; Niinemets et al. 2010; Monks et al. 2015). NMVOC emissions are a byproduct of photosynthesis, and also a response to various environmental factors, including temperature, atmospheric CO<sub>2</sub>, soil moisture, and drought stress. Land use changes can drastically affect isoprene (the dominant NMVOC) sources, for instance in the tropics where large scale deforestation has converted forest into crop lands (Kanakidou et al. 2005). This thesis focuses in this thesis I focus<sup>mh</sup> on emissions of isoprene in Australia.

Globally around 710 - 1150 Tg C yr<sup>-1</sup> of BVOCs are emitted (Guenther et al. 1995; Lathière et al. 2006; Guenther et al. 2012; Messina et al. 2016). BVOCs are emitted by vegetation, with the most dominant emitters being tropical trees. The main VOCs emitted are isoprene (C<sub>5</sub>H<sub>8</sub>) (~ 50 – 70%), and monoterpenes (C<sub>10</sub>H<sub>16</sub>) (~ 30%), which have different emission amounts depending on the tree species(Guenther et al. 2012; Sindelarova et al. 2014). Other emissions include methanol (CH<sub>3</sub>OH), ethanol (C<sub>2</sub>H<sub>6</sub>O), acetaldehyde (CH<sub>3</sub>CHO), acetone ((CH<sub>3</sub>)<sub>2</sub>CO), ethene (C<sub>2</sub>H<sub>4</sub>) and propene (C<sub>3</sub>H<sub>6</sub>) (Guenther et al. 2012). Many of these estimates come from Model of Emissions of Gases and Aerosols from Nature (MEGAN), a bottom-up biogenic emissions model.<sup>tb</sup> MEGAN parameterises BVOC emissions as a function of temperature, land cover, plant functional type, etc., with descriptions given in Guenther et al. (2012).<sup>tb</sup>

### 1.3.2 Biogenic emissions modelling

Many of the global estimates given in the paragraph above come from Model of Emissions of Gases and Aerosols from Nature (MEGAN), a bottom-up biogenic emissions model. Biogenic emissions models generally take bottom-up approach, where the environment is simulated and then the emissions from that environment are estimated based vegetation type, climate, and other local environmental factors. The most important parameters dealing with VOC emissions are leaf area index, emission factors, plant functional type, light availability, and temperature (Messina et al. 2016). The estimated emission factors are built up from measured plant emissions, which can be measured downwind (e.g., Nguyen et al. 2014) or from within a chamber (e.g., Paulot et al. 2009b; Nguyen et al. 2014). There are several models that estimate VOC emissions, such as MEGAN (created and published by (Guenther et al. 1995), with subsequent updates in Guenther et al. (2000), Guenther et al. (2006), and Guenther et al. (2012)), LPJ-GUESS (Lund-Potsdam-Jena General Ecosystem Simulator) (Arneth et al. 2007), and ORCHIDEE (Organising Carbon and Hydrology in Dynamic EcosystEm)

(Messina et al. 2016). Many of the available bottom-up models have inputs and sensitivities similar to those of MEGAN. MEGAN is used within this thesis as a module within a chemical transport model (Section 2.3.4).<sup>tb</sup>

The general formula governing modelled emissions  $E$  for a species  $i$  (from Brasseur and Jacob (2017)) is as follows:<sup>tb</sup>

$$E_i = A \times EF_i \times S_i \quad (1.5)$$

with  $A$  the activity rate,  $EF_i$  being the emission factors, and  $S_i$  is a scaling factor accounting for meteorology and other effects not included in  $A$  or  $F$  (e.g., seasonal temperature). For example, to estimate isoprene emission  $E_{isop}$  per second for an area, the equation becomes  $A$  = how many trees in an area multiplied by  $EF_{isop}$  = isoprene emitted per tree per second multiplied by  $S_{isop}$  = a scaling factor based on temperature and soil moisture. This is a simplified example of a bottom-up estimate for isoprene emissions.<sup>tb</sup>

There is a potential issue caused by shared underlying algorithms and assumptions between different emission models, which lead to an unrealistic convergence on isoprene emission estimation (Arneth et al. 2008). MEGAN can depend greatly on parameterised inputs such as temperature, sunlight, plant type, plant coverage, plant health, etc. (Arneth et al. 2008; Niinemets et al. 2010). Sensitivity to these factors is pervasive in bottom up emissions models (e.g., Marais et al. 2014; Miller et al. 2014; Messina et al. 2016). Relatively few Australian species are used within MEGAN's plant functional type parameterisations, and those that are may be poorly represented (Emmerson et al. 2016). Additionally there is uncertainty around the soil moisture parameterisations, with soil moisture and its impacts on BVOC emission poorly understood in Australia (Sindelarova et al. 2014; Emmerson et al. 2019). Modelled emissions are sensitive to soil moisture, especially near the wilting point, below which trees stop emitting isoprene and other VOCs completely as they can no longer draw water (Bauwens et al. 2016). This leads to increased uncertainty for Australian emissions.<sup>tb</sup>

The other common emission estimation technique is termed top-down. Top-down emissions estimates are derived from measurements of the child products of emitted VOCs. Based on top-down emissions estimates, isoprene emissions appear to be overestimated in Australia. Bauwens et al. (2016) showed that isoprene emissions were overestimated by up to a factor of 3 over Australia. Emmerson et al. (2016) suggest isoprene emission estimates are 2-6 times too high compared against available measurements of isoprene concentrations. They compared modelled data to campaign measurements from multiple sites over different seasons and found that scaling emissions did not universally improve model outputs.<sup>tb</sup>

### 1.3.3 Isoprene

Isoprene (2-methylbuta-1,3-diene) is the dominant BVOC emitted to the atmosphere. It is of major importance to the atmosphere, as it is involved in various processes that alter the oxidative capacity of the atmosphere. Isoprene affects  $\text{NO}_x$  and  $\text{HO}_x$  cycling, see for example reactions 1.1, 1.3. In the presence of  $\text{NO}_x$ , isoprene forms tropospheric ozone and SOA (Wagner 2002; Atkinson and Arey 2003; Millet et al. 2006; Patchen et al. 2007). It has a short lifetime during the day, roughly an hour due to OH oxidation

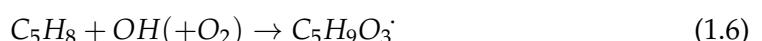
(Atkinson and Arey 2003). Ozonolysis and photolysis are lesser oxidation pathways, which can form similar products (Nguyen et al. 2016; Wolfe et al. 2016).

Measurements of isoprene are often uncertain, as the compound is reactive and short lived. Chamber experiments are used to determine how isoprene behaves once it is emitted into the atmosphere; however, laboratory derived reaction rates may be unsuitable when modelling the natural atmosphere, as natural air masses frequently differ from air masses used in chambers (Kanakidou et al. 2005; Nguyen et al. 2014). In the absence of extensive measurements, isoprene emissions and subsequent atmospheric oxidative chemistry are often modelled. MEGAN and other models(Guenther et al. 1995), and subsequent updates (Guenther et al. 2000; Guenther et al. 2006; Guenther et al. 2012),<sup>tb</sup> have been used throughout the atmospheric community to make global estimates of isoprene emissions, at roughly 500-600 Tg yr<sup>-1</sup>, emitted mostly during the day. Australia has increased uncertainty (relative to globally) due to its lack of measurements and study (see Section 1.3.2).<sup>tb</sup> Other models also exist which estimate similar isoprene amounts, such as LPJ-GUESS (Lund-Potsdam-Jena General Ecosystem Simulator) (Arneth et al. 2007), and ORCHIDEE (Organising Carbon and Hydrology in Dynamic Ecosystem) (Messina et al. 2016). There is a potential issue caused by shared underlying algorithms and assumptions between the models, which lead to an unrealistic convergence on isoprene emission estimation (Arneth et al. 2008). The global emission factors used to derive these estimates are based on modelling emissions from different plant species (phenotypes), and relatively few Australian species were used when forming these estimates. This leads to increased uncertainty for Australian emissions.<sup>tb</sup> Due to the highly reactive nature of isoprene, modelling of related chemical products is sensitive to uncertainties, for example the diurnal pattern of isoprene emissions affects modelled ground level ozone (Fan and Zhang 2004; Hewitt et al. 2011). The global uncertainty of isoprene emission was estimated to be a factor of 2 to 5 (250-750 Tg yr<sup>-1</sup>) (Kanakidou et al. 2005). Improvements over the years have been incremental, and generally localised to regions of particular interest for air quality such as China and the USA (Guenther et al. 2012; Jiang et al. 2018).

#### 1.3.4 Isoprene chemistry

Isoprene is emitted and enters the atmosphere in the gas phase, where it reacts quickly with OH and other radicals. One common compound that is produced by these reactions is HCHO, which can be measured from satellites and is often used to estimate how much isoprene is being emitted. Isoprene reacts with OH, ozone, or NO<sub>3</sub>, leading to organic peroxy radicals (ROO'). These go on to form many products and lead to (amongst other things) aerosol, formaldehyde, and ozone formation, depending on sunlight and NO<sub>x</sub> concentrations (Atkinson 2000).

Figure 1.5 shows the first stages of oxidation of isoprene by OH. The primary first step for atmospheric isoprene is photooxidation, reacting with OH to form isoprene hydroxyperoxy radicals (ISOPOO: C<sub>5</sub>H<sub>9</sub>O<sub>3</sub>, a subset of ROO'), and is shown in Equation 1.6 (Patchen et al. 2007; Wolfe et al. 2016; Marvin et al. 2017).



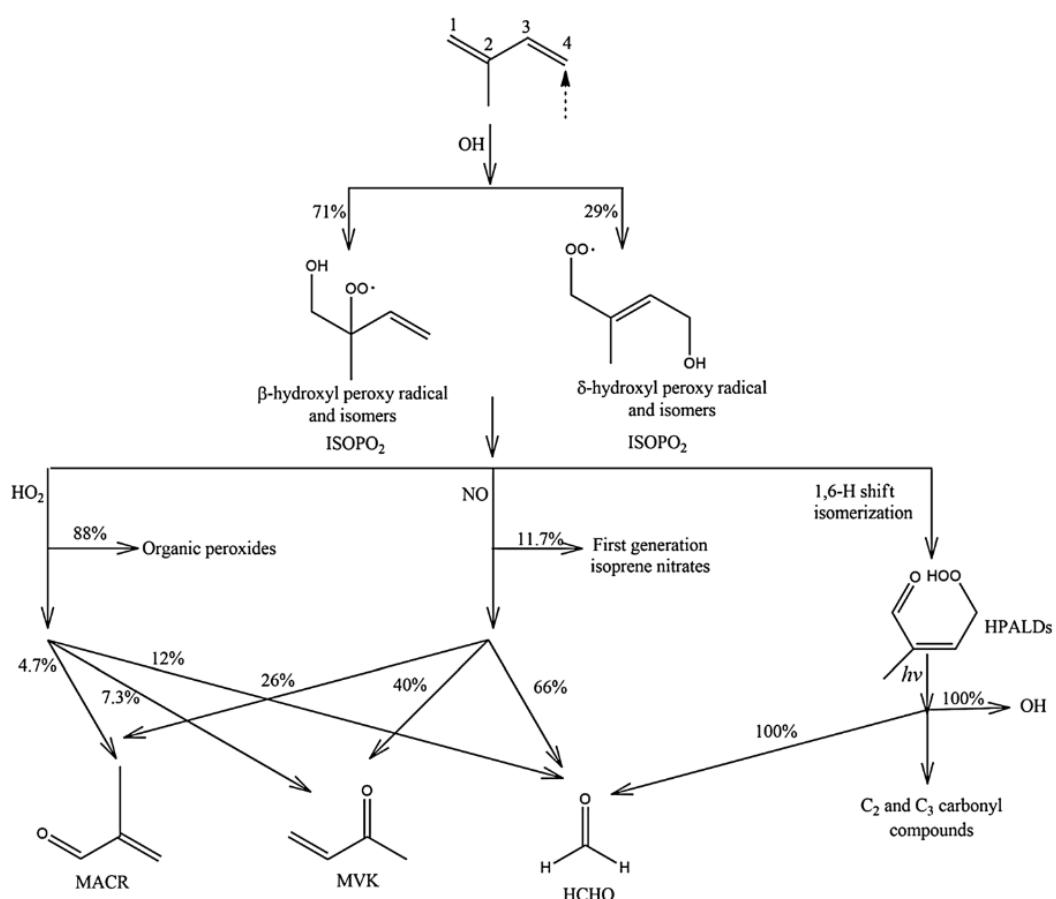
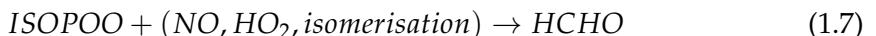


FIGURE 1.5: Simplified isoprene products following oxidation by OH, reproduced from Mao et al. (2013).

Different isomers are formed based on which carbon the OH adducts to. The subsequent processes that begin with isoprene oxidation are often called the isoprene (photochemical) cascade (e.g., Crounse et al. 2012; Paulot, Henze, and Wennberg 2012; Wolfe et al. 2016).

ISOPOO reacts with HO<sub>2</sub> or NO, producing stable products. This pathway also produces HCHO (with varying yields):



During the day HCHO has a lifetime of 1-2 hrs, while ISOPOO lasts ~ 100 s, making reaction 1.6 a rate limiting factor in HCHO production (Wolfe et al. 2016). Additionally, ISOPOO can isomerise and produce hydroperoxy-methyl-butenals (HPALDS) (see figure 1.5), which also leads to HCHO. At higher NO mixing ratios (at least a few hundred pptv), ISOPOO react mostly with NO. At low NO (less than 50 pptv), ISOPOO is more likely to either isomerise, react with HO<sub>2</sub>, or react with another ROO.

There is uncertainty about which pathways are most important following ISOPOO production, affecting predictions by atmospheric models (Nguyen et al. 2014). This limits understanding of the relative importance of some chemical processes, such as auto-oxidation of ISOPOO (Crounse et al. 2013). The reaction pathways depend on local concentrations of NO<sub>x</sub>: the high and low NO<sub>x</sub> pathways are dominated by NO and HO<sub>2</sub> reactions respectively. HO<sub>2</sub> reactions predominantly produce hydroxyhydroperoxides (ISOPOOH), while NO reactions produce isoprene nitrates (ISOPN) (Crounse et al. 2006). If measured, first generation ISOPN and ISOPOOH products can be used to determine the portion of isoprene oxidation following each pathway (e.g., Yu et al. 2016). Globally around one third of ISOPOO react with HO<sub>2</sub>, and two thirds react with NO (Paulot et al. 2009b). Most of these reaction pathways produce HCHO; however, the rate of production differs between different pathways, along with that of methyl vinyl ketone (MVK), and methacrolein (MACR), which can then be used to estimate reaction regimes (Marais et al. 2012; Liu et al. 2016b; Wolfe et al. 2016).

#### 1.3.4.1 High NO<sub>x</sub> pathway

In the presence of NO<sub>x</sub>, ISOPOO reacts with NO and forms ISOPN, which affect levels of both HO<sub>x</sub> (OH+HO<sub>2</sub>) and NO<sub>x</sub>. ISOPN generally act as a sink of HO<sub>x</sub>, and can be a sink or reservoir for NO<sub>x</sub> (Mao et al. 2013; Fisher et al. 2016). A portion of the ISOPN are recycled back to NO<sub>x</sub>, serving as a reservoir of nitrogen and allow its transport to the boundary layer of remote regions (Patchen et al. 2007; Paulot et al. 2009a). The nitrates can also build up in the winter, when removal processes are not as dominant (Lelieveld et al. 2009).

Under high NO<sub>x</sub> conditions there is a large and rapid yield of HCHO, with most of the ultimate HCHO production occurring within one day (Palmer et al. 2006). First generation ROO react with NO yielding MVK(~ 40%), MACR(~ 26%), and HCHO(~ 60%) which is higher than yields after reaction with OH (Liu et al. 2013; Mao et al. 2013). The MVK and MACR products form additional HCHO within a few hours due to oxidation by OH (Palmer et al. 2006).

### **1.3.4.2 Low NO<sub>x</sub> pathway**

In low NO<sub>x</sub> environments, ISOPOOH is formed in yields > 70%, while MACR, MVK, and HCHO are formed at ~ 5%, ~ 7%, and ~ 12% respectively (Paulot et al. 2009b; Mao et al. 2013). This ISOPOOH mostly reacts with OH to form dihydroxyperoxides (IEPOX) while regenerating OH (Mao et al. 2013). This pathway has lower and slower ultimate yields of HCHO from isoprene when compared to the high-NO<sub>x</sub> pathway (Palmer et al. 2006).

Isoprene oxidation and subsequent reactions are less well understood when lower concentrations of NO are present in the atmosphere. It was thought that in low NO environments, like those far from anthropogenic pollution and fires, oxidation of isoprene would create ISOPOOH and reduce local concentrations of OH and HO<sub>2</sub> (Guenther et al. 2000; Paulot et al. 2009b). However this reduction was not seen in measurements and HO<sub>x</sub> levels have been shown to be largely unaffected by isoprene concentrations (Paulot et al. 2009b). HO<sub>x</sub> is recycled through IEPOX, formed from ISOPOOH oxidation, and some HO<sub>x</sub> is produced in the formation of MACR and MVK (Paulot et al. 2009b). Though uncertain, OH and HO<sub>2</sub> production or regeneration mechanisms have improved model outcomes when compared against campaigns (Peeters and Muller 2010; Crounse et al. 2012).

Uncertainties and bias from measurements have made it more difficult to understand what happens in low NO<sub>x</sub> conditions, and many observations of OH remain under-predicted in models (Mao et al. 2012). Due to OVOC interference, measurements in low NO<sub>x</sub> environments can lead to massively overestimated MVK and MACR yields (Nguyen et al. 2014). Nguyen et al. (2014) show preliminary estimates of low-NO yields of MVK and MACR to be 6±3% and 4±2% respectively, consistent with Liu et al. (2013), but only when cold-trapping methods are employed. Many instruments have been shown to generate OH internally, creating anomalous VOC readings due to within-instrument oxidation (Mao et al. 2012).

Improved understanding of both the chemistry and instrument sensitivities has helped closed the gap between model predictions and detected concentrations of VOCs and OH (Mao et al. 2012), but uncertainties remain in isoprene oxidation mechanisms. Examples (taken from Nguyen et al. (2014)) include:

- isoprene nitrate yields, which range from 4-15% (Fisher et al. 2016)
- 90% disagreements in MACR and MVK yields (Liu et al. 2013)
- unknown HPALD fates (Peeters, Nguyen, and Vereecken 2009; Crounse et al. 2013)
- poorly-characterised RO<sub>2</sub> lifetime (Wolfe et al. 2012).

### **1.3.4.3 Night time processes**

At night when OH concentrations have dropped, isoprene can remain in the atmosphere. Typically less than half of this night time isoprene is removed through ozonolysis (Atkinson and Arey 2003). A build up of NO<sub>3</sub> radicals can be seen at night, when photolysis is not removing them (Atkinson 2000; Brown et al. 2009). This build up is

enhanced in polluted (high  $\text{NO}_x$ ) areas.  $\text{NO}_3$  are largely formed through ozone reactions. In these conditions isoprene is consumed by  $\text{NO}_3$  radicals that join to one of the double bonds, producing organic nitrates ( $\text{RONO}_2$ ) in high yield (65% to 85%) (Mao et al. 2013).

In areas with high  $\text{NO}_x$  levels, greater than 20% of the isoprene emitted late in the day ends up being oxidised by the  $\text{NO}_3$  radical overnight (Brown et al. 2009). At night isoprene affects both  $\text{NO}_x$  concentrations and ozone levels, and can form harmful organic nitrates and SOA (Brown et al. 2009; Mao et al. 2013). These nitrates go on to produce further SOA, largely due to  $\text{NO}_3$  reacting with first generation isoprene oxidation products (Rollins et al. 2009). The night-time concentrations of OH and ozone also have a complex effect on  $\text{NO}_x$  removal in high latitude winters, when photolysis and NO reactions are reduced (Ayers and Simpson 2006).

### 1.3.5 Radiative Forcing

Another reason for interest in VOCs is that they are a dominant source of organic aerosol, which are a major source of uncertainty in radiative forcing. VOC emissions affect ozone along with several atmospheric parameters that directly and indirectly alter radiative forcing rates (e.g., Arneth et al. 2008). This is even more important in Australia where VOCs are so poorly represented by contemporary modelling (Emmerson et al. 2016). Aerosols are particles in the atmosphere, and they affect radiative forcing in ways that are difficult to accurately quantify. For some years it has been understood that aerosols have an overall cooling effect within the atmosphere. Smaller particles can match the wavelengths of visible light, interfering with incoming radiation (Kanakidou et al. 2005). Aerosol products from gas phase emissions (or the children thereof) also play an indirect and complex role in cloud properties, with a net cooling effect (Kanakidou et al. (2005), Stocker et al. (IPCC, 2013: *Climate Change 2013: The Physical Science Basis. Contribution of Working Group I to the Fifth Assessment Report of the Intergovernmental Panel on Climate Change*, Chapter 7, 8)). Aerosol and cloud formation remain as large uncertainties in recent IPCC reports (Forster et al. 2007). Figure 1.6 shows that uncertainty from aerosol effects dominate the uncertainty in radiative forcing.

## 1.4 Formaldehyde

In conjunction with atmospheric chemistry and radiative models, satellite measurements quantify the abundance of HCHO in the atmosphere. Isoprene is hard to measure directly due to its short lifetime and weak spectral absorption, instead HCHO is often used as a proxy (Millet et al. 2006; Fu et al. 2007; Dufour et al. 2008; Marais et al. 2012; Bauwens et al. 2013; Kefauver, Filella, and Peñuelas 2014; Bauwens et al. 2016; Surl, Palmer, and Abad 2018). The existence of satellite data covering remote areas provides an opportunity to improve VOC emissions estimates leading to more robust models of global climate and chemistry. This method is implemented within this thesis (Section 3.2), and relevant information dealing with HCHO is provided here.

Formaldehyde (HCHO), also known as methanal, methyl aldehyde, or methylene oxide, is part of the aldehyde family. HCHO is an oxygenated VOC that is toxic,

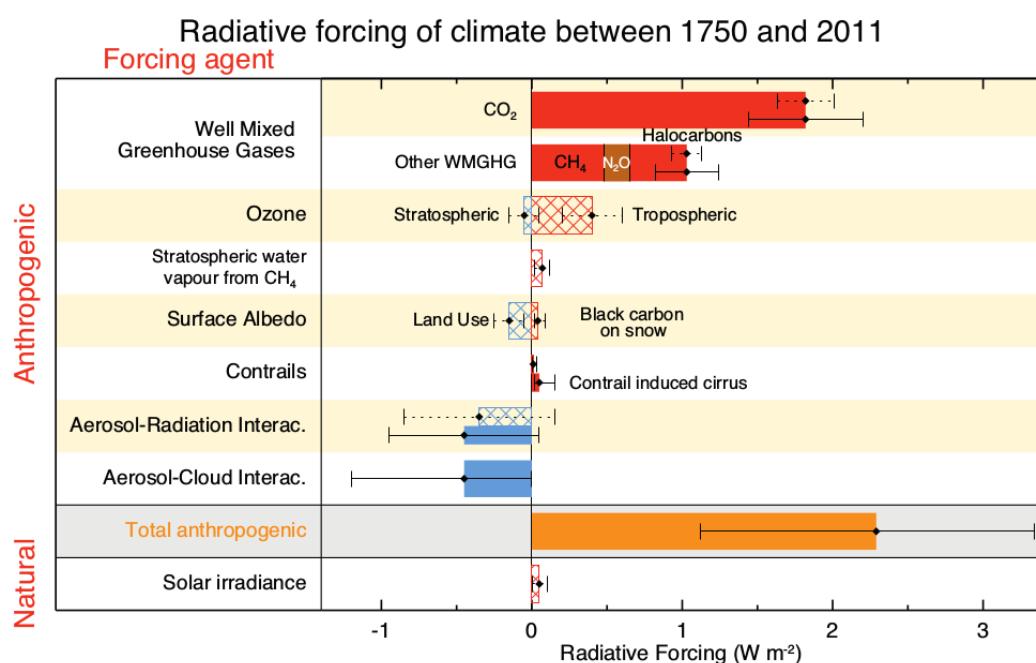
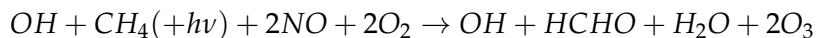


FIGURE 1.6: The overall radiative forcings and uncertainties of several atmospheric constituents. Reproduced from Stocker et al. (IPCC, 2013: *Climate Change 2013: The Physical Science Basis. Contribution of Working Group I to the Fifth Assessment Report of the Intergovernmental Panel on Climate Change*), chapter 8.

allergenic, and a potential carcinogen. Generally, HCHO is not dangerous at low background atmospheric levels. Globally, HCHO production is mainly due to the oxidation of methane; however, over continental regions, HCHO enhancement above the background level is largely due to isoprene emissions. HCHO production also depends on NO<sub>x</sub> concentrations, which affect the yield from isoprene oxidation. Isoprene reactions yield more HCHO in the high-NO<sub>x</sub> pathway (compared to the low-NO<sub>x</sub> pathway) (Marais et al. 2012). HCHO measurements are often used to evaluate how well isoprene chemistry is simulated by models, as HCHO levels depend on initial VOCs and oxidants (Marvin et al. 2017).

#### 1.4.1 Sources and sinks

Background levels of HCHO in the atmosphere are driven by the oxidation of CH<sub>4</sub> by the OH, which produces  $\sim 970 \text{ Tg yr}^{-1}$  (Fortems-Cheiney et al. 2012). Atkinson (2000) summarised the background formation of HCHO with the following reaction:



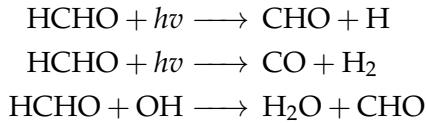
This shows that photolysis and oxidation of methane forms HCHO and ozone in a process that regenerates the OH radicals. CH<sub>4</sub> concentrations are relatively well constrained in models, with the Atmospheric Chemistry and Climate Model Inter-comparison Project showing only  $\sim 3\%$  inter-quartile range (Young et al. 2013).

Within the continental boundary layer, HCHO is enhanced above background levels due to NMVOC emissions reacting with OH radicals in the presence of NO<sub>x</sub> (Wagner 2002; Millet et al. 2006; Kefauver, Filella, and Peñuelas 2014). The total contribution from NMVOC oxidation is  $\sim 358 \text{ Tg yr}^{-1}$  (Fortems-Cheiney et al. 2012). Enhancements to regional and continental HCHO are largely driven by emissions of isoprene (Guenther et al. 1995; Palmer et al. 2003; Shim et al. 2005; Kefauver, Filella, and Peñuelas 2014). The exception is where emissions of HCHO or precursors occur due to fires or anthropogenic activities such as fossil fuel combustion, natural gas flaring, ethanol refining, and agricultural activity (Guenther et al. 1995; Kefauver, Filella, and Peñuelas 2014; Wolfe et al. 2016). Other terpenoids (monoterpenes, sesquiterpenes, etc.) can also produce HCHO, although generally to a lesser extent than isoprene, methane and biomass burning (Guenther et al. 2012). Anthropogenic sources of HCHO are largely negligible; however, their signals can be seen in very large cities or near large industrial sources (Millet et al. 2008; Zhu et al. 2014b).

In the past, HCHO levels were underestimated by models, often with large discrepancies, due to the poor understanding of methyl peroxy radical (CH<sub>3</sub>OO) chemistry (Wagner 2002). Now HCHO concentrations are better understood, however precursor emissions are one of the main unknowns (e.g., Emmerson et al. 2016; Marvin et al. 2017). Another source of discrepancy between modelled and measured HCHO concentrations is second and later generational isoprene oxidation chemistry (Marvin et al. 2017).

HCHO has two major sinks totalling  $\sim 1210 \text{ Tg yr}^{-1}$ , reactions with OH (oxidation), and photolysis (Levy 1972; Crutzen, Lawrence, and Poschl 1999; Wagner 2002; Fortems-Cheiney et al. 2012; Kefauver, Filella, and Peñuelas 2014). The other sinks are

wet and dry deposition, although these are not as significant ( $\sim 32 \text{ Tg yr}^{-1}$ ) (Atkinson 2000; Fortems-Cheiney et al. 2012). Oxidation and photolysis reactions begin as follows (Ayers et al. 1997):



These reactions lead to a daytime lifetime of a few hours (Atkinson 2000; Millet et al. 2006). Both these loss processes (photolysis, oxidation) form CO, and lead to production of hydroperoxyl radicals ( $\text{HO}_2$ ). These products have global significance to radiative forcing and oxidative capacity (Franco et al. 2015).

#### 1.4.2 Measurement techniques

HCHO measurements are used extensively throughout this thesis, and a brief introduction to the relevant techniques is provided here. HCHO data used in this thesis have been measured using two different techniques: Fourier Transform Infra-Red (FTIR) Spectrometry and Differential Optical Absorption Spectroscopy (DOAS). FTIR uses the Fourier transform of a measured spectrum to quantify species which interfere with the spectrum in the infra-red region. DOAS methods are based on light interference and absorption through air masses.

The DOAS technique takes advantage of the optically thin nature of HCHO in order to linearise the radiance differential through air masses with and without HCHO, using the Beer-Lambert intensity law. This method is used globally both from ground-based and space-based instruments for HCHO detection (Guenther et al. 1995; Gonzalez Abad et al. 2015; Davenport et al. 2015). As a trace gas, HCHO absorbs light over a few wavelength bands, which allows instruments to detect concentrations between a known light source and a detector. Figure 1.7 shows the interference spectrum of HCHO along with a typical band used to examine interference in the DOAS technique. FTIR and DOAS measurements have a range of uncertainties, including systematic and random measurement errors and uncertain a priori shape factors and water vapour profiles (e.g., Franco et al. 2015). One difficulty is that this interference is relatively small (HCHO is optically thin) and other compounds absorb light at similar wavelengths (Davenport et al. 2015).

Other types of measurement involve directly measuring the air, and determining chemical compounds through their physical properties such as by mass spectrometry analysis of mass to charge ratios ( $m/z$ ) of ionised air masses. Two examples of this include proton transfer reaction mass spectrometers (PTR-MS), and gas chromatography mass spectrometers (GC-MS). These instruments can be also used to determine the gas phase evolution of other isoprene and monoterpene products (e.g., Lee et al. 2006; Nguyen et al. 2014; Wolfe et al. 2016; Lerner et al. 2017). Reasonable agreements between different instruments and techniques can be achieved, although titration and calibration differences can lead to large ( $\sim 30\%$ ) discrepancies (e.g., Hak et al. 2005). These differences and non-uniformity between measurements (even among identical

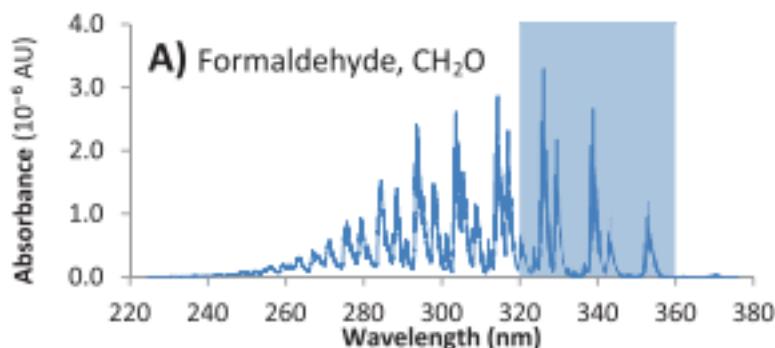


FIGURE 1.7: HCHO spectrum, with a typical band of wavelengths used for DOAS path measurements. Reproduced from Davenport et al. (2015).

instruments) are part of the reason HCHO does not have a consistent network for global measurements like those for greenhouse gases or ozone (Fortems-Cheiney et al. 2012).

#### 1.4.2.1 Satellite measurements

Satellites remotely sense atmospheric HCHO through irradiance measurements of solar light that has reflected off the earth's surface. These irradiances are affected by gases that absorb radiation along the reflected path of light between the detector, earth, and sun. The irradiance is then used to estimate how much of a particular gas exists along this path, which gives an estimate called the slant column. The retrieved slant column of a particular gas (species) can be transformed into a vertical column by scaling the path length in conjunction with accounting for the light scattering properties of the trace gas. The scaling coefficient created to transform from slant to vertical column is called the Air Mass Factor.

Several satellites provide long term HCHO observations with near complete global coverage. Some of these are:

- ERS-2 launched in April 1995, housing the GOME ultraviolet and visible (UV-Vis) spectrometer
- AURA launched in July 2004, housing the OMI UV-Vis spectrometer
- MetOp-A and B launched in October 2006 and September 2012 respectively, both housing a GOME-2 UV-Vis spectrometer.

These satellites are on low earth orbit trajectories and overpass any area up to once per day. Satellites use DOAS techniques (Section 1.4.2.2) with radiative transfer calculations on solar radiation absorption spectra to measure column HCHO. An example of a spectrum retrieved from the GOME-2 instrument is given in figure 1.8.

Satellite observations often require both remote-sensing and in situ measurements combined with modelled data for validation (Marais et al. 2014). There is less information available from satellite measurements at higher latitudes due to increased error in measurements over the more slanted column paths (De Smedt et al. 2015). Validation

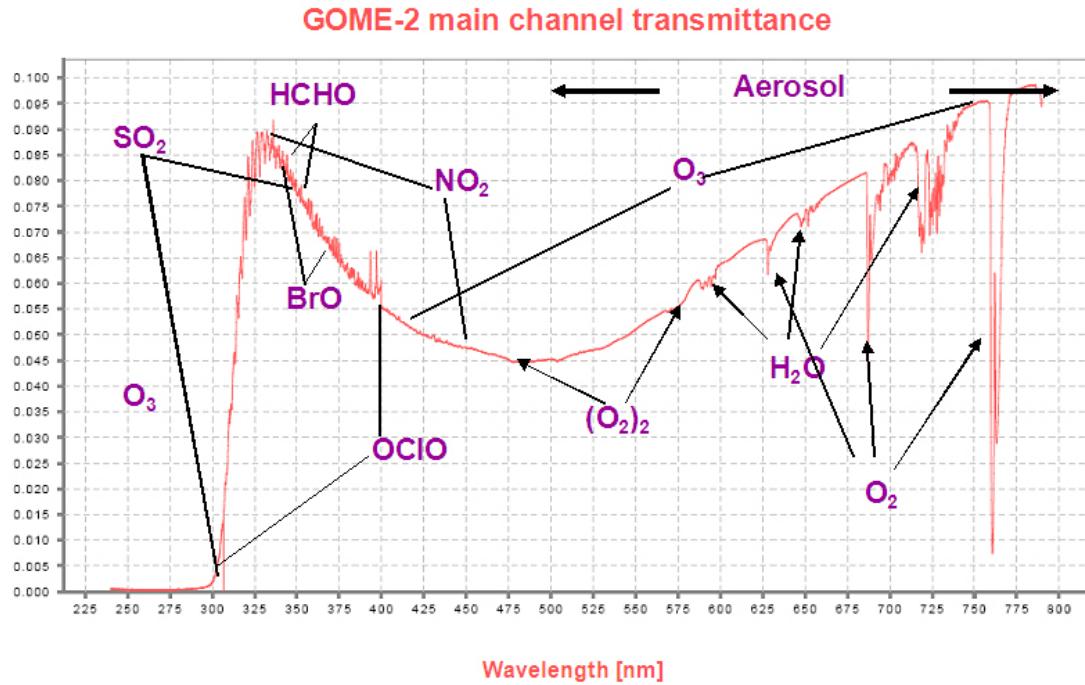


FIGURE 1.8: An example spectrum showing absorption features used for species concentration measurements by GOME-2. Image reproduced from EUMETSAT and ESA (EUMETSAT 2015).

is important due to the various uncertainties in the satellite remote sensing process. This can be done using aircraft data to provide model validation (e.g., Zhu et al. 2016). Satellites use an assumed shape factor to improve the vertical column estimate; however, this can lead to a bias between satellite data and other measurements (Zhu et al. 2016). Different satellites, instruments, or techniques that measure HCHO (or any trace gas) can give different results due to differing a priori assumptions (Lorente et al. 2017). The concept of differences that arise between datasets based on measurement techniques or underlying instrument biases is called structural uncertainty.

#### 1.4.2.2 DOAS

The DOAS technique uses solar radiation absorption spectra to measure trace gases through paths of light. Beer's law states

$$T = I / I_0 = e^{-\tau} \quad (1.8)$$

with  $T$  being transmittance,  $\tau$  being optical depth, and  $I, I_0$  being radiant flux received at instrument and emitted at source respectively. The Beer-Lambert law of extinction allows spectroscopic measurement of absorbing chemical species (absorbers) in the atmosphere:

$$I_B = I_{B_0} e^{-\tau_s} \quad (1.9)$$

where  $I_B$ ,  $I_{B_0}$  is backscattered intensity with and without the absorber respectively, and  $\tau_s$  is the optical thickness of the absorber along the measured path between source and instrument.

$\tau$  can be described using the scattering and absorption cross section area ( $\alpha$ ,  $\text{cm}^2$ ) and density ( $\eta$ , molec  $\text{cm}^{-3}$ ) of an absorber as follows:

$$\tau = \int \alpha(s) \eta(s) ds \quad (1.10)$$

Here  $\tau$  represents the sum of optical thicknesses of each absorber within the measured path ( $s$ ) through a medium. Substituting Equation 1.10 into Equation 1.9 leads to

$$I = I_0 \exp \left( \sum_i \int \eta_i \alpha_i ds \right) \quad (1.11)$$

Where  $i$  represents a chemical species index, and the integral over  $ds$  represents integration over the path from light source to instrument.

Another way of describing optical depth (also called optical thickness) is the natural logarithm of the ratio of incident radiant power to transmitted radiant power through a material (from Equation 1.9). In the atmosphere, ~~the optical depth of various chemical species can be determined using incoming solar radiation. we are interested in the optical depth of various chemical species, and we use incoming solar radiation to determine this.~~<sup>mh</sup> The difference between solar radiation at the top of the atmosphere and the earth's surface defines the atmospheric optical depth along the path of observation.

$$\tau = \ln \frac{\phi_e^i}{\phi_e^t} \quad (1.12)$$

where  $\phi_e^i$  is radiant flux at the surface,  $\phi_e^t$  is the solar radiant flux that arrives at the top of the atmosphere. In the atmosphere, optical depth can be due to several factors including scattering, chemical absorbance, and aerosols.

## 1.5 Atmospheric Chemistry Modelling

Models can fill the gaps (both spatial and temporal) in measurement records, and ~~can help us~~<sup>mh</sup> improve ~~our~~<sup>mh</sup> understanding of the natural world. For example, they can be used to examine future outcomes resulting from changing ~~our~~<sup>mh</sup> emissions. They can be used to increase measurement accuracy (for instance in satellite measurements) and determine where ~~information is lacking~~<sup>we lack information</sup><sup>mh</sup>, while also evaluating the performance of new instruments. Some advanced measurement techniques make use of modelled a priori information in order to produce useful outputs. Models of ozone in the atmosphere are used broadly for international assessments of ozone precursor emissions, and estimating effects from related processes (such as radiation) (Young et al. 2017). Models provide an estimate of many trace gas concentrations; however, verification is required, and is generally performed using results from measurement campaigns. In situ measurements from campaigns or measurement stations can be used to examine what is happening at a particular location. These data are

used to determine how accurate models or estimates are - however the utility is limited to where and when the measurements took place.<sup>tb</sup> Precisely representing various chemicals and reactions in the atmosphere allows efficient mitigation of pollution, since proposed scenarios can be compared against one another. Currently, models require improved isoprene emissions and subsequent chemistry for effective air quality determination (Marvin et al. 2017).

### 1.5.1 Box models

Box models simulate chemistry in a singular set of conditions without transport or spatial gradients. These models often parameterise things such as transport and emissions that would realistically take place at the edges of the box. Box models can be used to test chemical mechanisms in specific scenarios, such as high or low NO<sub>x</sub> environments. For example, Marvin et al. (2017) used a box model matching conditions in southeast USA to evaluate isoprene mechanisms from several models.

By allowing for interactions between boxes this concept can be extended to multiple-box models. Multiple-box models are simply multiple instances of single boxes with the addition of transport between them. Transport requires meteorological fields such as wind velocities and turbulence. The meteorology fields can be modelled, and/or input as parameters.

### 1.5.2 Chemical transport models

Chemical transport models (CTMs) provide a simulation of chemical densities and transport over time, through the atmosphere. Chemistry in the atmosphere is a complex system of coupled reactions and dynamics, which can be solved using numerical partial differential equation solvers. Chemical models require many inputs such as meteorological conditions and emissions. Initial (atmospheric starting state) and boundary (inputs and outputs at the edge of the modelled system) conditions are required, and models or inventories of emissions often make up inputs at the surface of the atmosphere.

CTMs simulate production, loss, and transport of chemical species. This is generally calculated using one or both of the Eulerian (box) or Lagrangian (puff) frames of reference. Eulerian models use equations of chemistry within and transport between volumes in a gridded spatial coordinate system, while Lagrangian models evaluate behaviour within a potentially changing frame of reference (for example within a cloud). CTMs normally solve continuity equations simultaneously for many coupled species. The continuity equations describe transport of a conserved quantity such as mass or energy, which, solved together with production and loss of a chemical can provide detailed simulations of natural processes.

The general continuity equation links a quantity of a substance ( $q$ ) to the field in which it flows and can be described by the formula:

$$\frac{\partial \rho}{\partial t} + \nabla \cdot j = \sigma$$

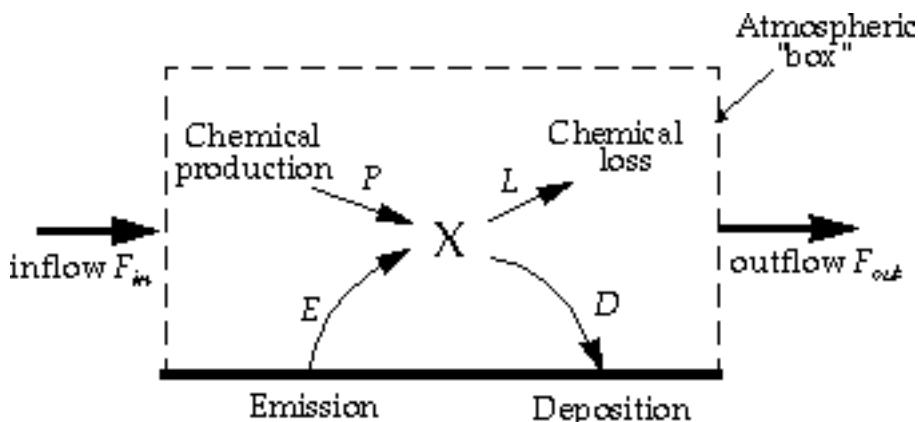


FIGURE 1.9: Standard box model parameters, reproduced from Jacob (1999).

where  $\rho$  is density of  $q$  in the field,  $t$  is time,  $\nabla$  is divergence,  $j$  is the flux ( $q$  per unit area per unit time entering or leaving the field), and  $\sigma$  is the generation or loss of  $q$  per unit volume per unit time.

The type of model best suited to modelling the entire earth uses the Eulerian frame of reference, where the atmosphere is broken up into 3-D boxes with densities and transport calculated and stored for sequential steps in time at each location. The mass balance equation must be satisfied in any realistic long term model and is as follows:

$$\begin{aligned}\frac{dm}{dt} &= \sum \text{sources} - \sum \text{sinks} \\ &= F_{in} + E + P - F_{out} - L - D\end{aligned}$$

where  $m$  is mass of a chemical,  $E$  and  $D$  are emission and deposition,  $P$  and  $L$  are production and loss, and  $F$  is chemical transport in and out, as shown in figure 1.9. Any large chemical model will solve this mass balance equation over highly coupled arrays of partial differential equations, which becomes computationally expensive as complexity increases.

Contemporary models generally use mathematical differential solving tools of various complexity (often called chemical mechanisms) to solve chemical equations in order to predict chemical species evolution over time. A chemical mechanism is a closed system of chemical reactions and their associated rate constants. Simplifications are required due to the large number of reactions that occur in the atmosphere, and the coupled and stiff nature of these reactions that slow down computation of the solutions (Brasseur and Jacob 2017). Mathematical stiffness in chemical systems of differential equations is due to the massively differing reaction time scales - for instance hydroxyl radicals react within seconds while methane has an atmospheric lifetime of 8-10 years (Wuebbles and Hayhoe 2002).<sup>tb</sup> Different solvers may be slower or faster, and some are more suited to particular situations. The choice of chemical mechanism is normally driven by the mathematical properties of the equations and systems being modelled (such as stability and stiffness), as well as limitations around processing power. Grouping together subsets of model equations or chemical species is often

performed and it is possible to use different solvers on each grouping. For example: since  $[O] \ll [O_3]$  the chemical family  $O_X$  ( $O_X \equiv O + O_3$ ) can be used to simplify chemistry simulations and approximate  $O_3$  concentrations (Brasseur and Jacob 2017, Chapter 3). Different chemical mechanisms may find different solutions to the same problems, due to how the numerical solvers are implemented, which can affect model output (Zhang et al. 2012).

The major chemical transport model used in this thesis is GEOS-Chem, an Eulerian 3-D box model as discussed above. In part, GEOS-Chem chemistry mechanisms are based on atmospheric chemistry chamber studies. In a chamber with clean air and high NO concentrations, isoprene photooxidation is initially driven by OH addition, followed by  $NO_x$  chemistry (150 min - 600 min), and finally  $HO_x$  dominated chemistry. Formation of isoprene nitrates (ISOPN) affect ozone levels through  $NO_x$  sequestration, the yields and fate of these nitrates was analysed in Paulot et al. (2009a). Prior to 2012, oxidation chamber studies were performed in high NO or  $HO_2$  concentrations, giving peroxy lifetimes of less than 0.1 s (Crounse et al. 2012; Wolfe et al. 2012). In most environments NO and  $HO_2$  concentrations are not so high. Crounse et al. (2011) examined the isomerisations associated with the oxidation of isoprene to six different isomers of  $ROO$  formed in the presence of oxygen.<sup>tb</sup>

~~Section 1.5.3 Emissions has been moved from here into new Section 1.3.2<sup>tb</sup>~~

~~The general formula governing modelled emissions  $E$  for a species  $i$  (from Brasseur and Jacob (2017)) is as follows ...<sup>tb</sup>~~

### 1.5.3 Uncertainties

This section summarises the major uncertainties models have in relation to VOCs, and ozone. Atmospheric chemical models by necessity use simplifications of real world processes, and also utilise information that may be itself uncertain or extrapolated. Uncertainty is introduced through both of these channels as well as through computational limitations.

#### 1.5.3.1 Emissions Inventories

Model results can be greatly affected by the choice of emissions inventory used to provide boundary conditions. Natural (biogenic or pyrogenic) and anthropogenic emissions often drive a large fraction of atmospheric oxidation and radical chemistry, especially in the continental boundary layer. Emissions inventories have been found to be reliable at larger (regional to global) scales, as long as they are derived from accurate input measurements (Zeng et al. 2015). Modelled ozone concentrations have been found to be sensitive to isoprene emissions and  $NO_x$  sources, both of which are uncertain within approximately a factor of 2 (Christian, Brune, and Mao 2017). Bottom up inventories of BVOCs (Section 1.3.2)<sup>tb</sup> remain largely uncertain due to missing or extrapolated plant functional type information, changing land cover, and parameterised environmental stressors (Guenther et al. 2000; Kanakidou et al. 2005; Millet et al. 2006). Global emissions inventories like MEGAN often implement data extrapolation, which also introduces uncertainties (Miller et al. 2014).

Modelled surface ozone may be overestimated if  $NO_x$  emissions are too high, due to their affects on the oxidative capacity of the atmosphere (Travis et al. 2016).  $NO_x$

and isoprene emissions have been shown to be the most significant sources of uncertainty for ozone concentrations near the surface over the US, while isoprene-derived products and lightning NO<sub>x</sub> drives uncertainty in the upper atmosphere (Christian, Brune, and Mao 2017).

Estimates of isoprene emission are based on a few algorithms that ~~may be heavily influenced by parameterised inputs (see Section 1.3.2). can depend greatly on parameterised inputs such as temperature, sunlight, plant type, plant coverage, plant health, etc. (Arneth et al. 2008; Niinemets et al. 2010).~~<sup>tb</sup> Sensitivity to these factors is pervasive in bottom up emissions models (e.g., Marais et al. 2014; Miller et al. 2014; Messina et al. 2016). Arneth et al. (2008) argue that the monopoly of isoprene emission estimates may be leading ~~us~~<sup>mh</sup> to an incorrect understanding of isoprene chemistry. VOC emissions form an integral part of many chemistry models, and poor characterisation of these can lead to many more problems in other models (e.g., Yue, Unger, and Zheng 2015). Models that depend on VOC emissions acquire the uncertainties and sensitivities (for example, to light and temperature parameters) found in VOC emissions (Yue, Unger, and Zheng 2015). This can be mitigated where measurements are used to constrain parameters (e.g., Stavrakou et al. 2014). ~~Modelled emissions are sensitive to soil moisture, especially near the wilting point, below which trees stop emitting isoprene and other VOCs completely as they can no longer draw water (Bauwens et al. 2016).~~ MEGAN accounts for soil moisture through a parameterisation that drops plant emissions to zero below a prescribed soil moisture level (the wilting point).<sup>tb</sup> For example,<sup>tb</sup> Jiang et al. (2018) found that improving MEGAN~~the~~<sup>tb</sup> parameterisation of drought based on a measurement campaign in the U.S. would lower isoprene emissions globally by  $\sim 17\%$ .

Chamber studies are often used to inform modelled isoprene reactions (e.g., Paulot et al. 2009b). However these results come with their own uncertainties, not the least of which comes from the difficulty of reproducing ambient air in laboratory conditions (Nguyen et al. 2014). Structural uncertainties (differences between different measurement techniques) in measurements also occur, which increases the difficulty of assessing isoprene datasets.

### 1.5.3.2 Resolution

Atmospheric chemistry simulations are somewhat sensitive to the gridbox resolution. Reducing model resolution can increase OH concentrations and ozone production rates (Wild and Prather 2006). Increasing model resolution can also impact OH, HO<sub>2</sub>, and ozone concentrations (e.g., Christian, Brune, and Mao 2017). Yu et al. (2016) show that only at higher resolution (0.25 by 0.3125°) does isoprene oxidise under the correct NO<sub>x</sub> pathway (through high or low NO<sub>x</sub> pathways, see Section 1.3.4) in variable NO<sub>x</sub> environments. This leads to an increase of high NO<sub>x</sub> pathway oxidation of isoprene at the lower resolutions, which leads to an overestimation of HCHO but not ozone at coarser resolutions. However, for many global scale analyses, errors from resolution are less important than those from chemistry, meteorology, and emissions (Christian, Brune, and Mao 2017; Christian et al. 2018).

### 1.5.3.3 Chemistry mechanisms

Chemical mechanisms in several contemporary models are likely inadequate (Marvin et al. 2017), especially for isoprene oxidation, with changes to reaction rates being inadequate to fix discrepancies from measurements. Ozone uncertainties are also affected by inadequate mechanisms, for example, ozone concentrations simulated by GEOS-Chem are most sensitive to NO<sub>2</sub> photolysis, the NO<sub>2</sub> + OH reaction rate, and precursor emissions (Christian, Brune, and Mao 2017).

Until recently, models struggled to represent OH in the atmosphere, due to missing chemistry and issues in measurement techniques. Prior to Mao et al. (2012), measurements of OH in high VOC regions may have been up to double the real atmospheric OH levels, due to formation of OH inside measuring instruments. Photolysis of photolabile peroxy-acid-aldehydes generates OH, improving model agreement with continental observations. OH and HPALD interactions are central to maintaining the OH levels in pristine and moderately polluted environments, which makes isoprene both a source and sink of OH (Peeters and Muller 2010; Taraborrelli et al. 2012). Mao et al. (2013) showed that drastically lowering (by a factor of 50) the rate constant for ROO isomerisation led to better organic nitrate agreements with measurements. These chemical updates led to more accurate modelling of OH concentrations, especially in low NO<sub>x</sub> conditions common in remote forests.<sup>1b</sup>

### 1.5.3.4 Clouds

One of the major uncertainties in chemical models is cloud formation and dynamics. Clouds are remarkably complex at a much finer scale than can be accurately modelled by global chemistry models (with current processing power). Globally over half (50-60%) of the world is covered by clouds, with ~ 10% of them being rain-clouds (Kanakidou et al. 2005). Wet scavenging in clouds not only depends on large scale cloud processes, but also on the microphysics of aerosols being scavenged, and these processes must be parameterised in coarse resolution models.

## 1.6 Australia and the southern hemisphere

Australia is unique, with its own climate, soil moisture, clay content, plant species, ecosystem life-cycles, agricultural and mining practices, transport corridors, and many other important properties that affect emissions, atmospheric chemistry, and ultimately atmospheric composition. Many regions in Australia are difficult or expensive to reach, and measurement campaigns are limited. In Australia, most long term air quality or composition measurements are performed in or near large cities. Australia is dominated by areas with little anthropogenic influence, and few ground based measurements of natural emissions take place (VanDerA et al. 2008). Since many Australian cities are on the edge of regions with rich VOC emissions, it is very important to clarify the quantity, type, and cause of VOC emissions. Understanding of emissions from these areas is necessary to inform national policy on air pollution levels.

Ozone enhancements above the background levels are most sensitive to emissions (of precursor gases), with meteorology, and atmospheric composition also important. Anthropogenic emissions of ozone precursors are important but relatively stable, while pyrogenic sources are greatly variable and dependent on weather, fuel, and fire intensity (e.g., Lawson et al. 2017). Biomass burning in southern Africa and South America has previously been shown to have a major influence on atmospheric composition in Australia (Oltmans et al. 2001; Gloudemans et al. 2006; Edwards et al. 2006), particularly from July to December (Pak et al. 2003; Liu et al. 2016a). Local fires are even more influential and the burning season for Australia can be all year, although distinct burning seasons for different regions are also apparent (Russell-Smith et al. 2007). Bushfire severity depends on regional vegetation, recent and current weather, and El Niño phase.

### 1.6.1 Ozone

Surface ozone levels over Australia are relatively low ( $\sim 20$  ppb) (Young et al. 2017). It remains unclear how ozone will change in the future as relatively little is known about precursors and influx for the continent. Australian air quality is monitored independently within each state. Measurement stations are generally located in population centres, which means there is a lack of natural emissions. Natural emissions can be important in urban regions as they can characterise the inflow, and affect air quality through production of  $O_3$  and other pollutants. Overall in the southern hemisphere there are relatively few records of ozone (Huang et al. 2018). This affects the scientific community's our<sup>mh</sup> ability to accurately determine sources of ozone in the troposphere, with current southern hemisphere trends lacking full explanation (Zeng et al. 2017).

Generally STT of ozone over Australia only affects the upper troposphere; however, ozone enhancements can reach low altitudes during heavy storms and cyclonic weather patterns (Alexander et al. 2013). The contribution of STT to overall tropospheric ozone budgets remains uncertain, especially in the southern hemisphere (Škerlak, Sprenger, and Wernli 2014). STT can enhance surface ozone concentrations above legal air quality limits (e.g., Lelieveld et al. 2009; Lin et al. 2015). Detecting ozone enhancements over the background profile in the relatively clean southern ocean atmosphere is simple. However, measurements of ozone over this region are sparse (Škerlak, Sprenger, and Wernli 2014), and quantification of transported ozone is difficult without large scale extrapolations. Ozone enhancements over the southern ocean signify either transported pollution or stratospheric influx (Jacobson and Hansson 2000). Quantifying ozone processes over the southern ocean could improve our<sup>mh</sup> understanding of chemistry in the "clean background environment", while additionally helping to validate model and satellite datasets. This poorly understood background composition forms the inflow of air into urban areas, and an improved understanding is required in order to model and attribute pollution and composition in these areas also.

## 1.6.2 Biogenic VOCs

The vegetation in Australia is diverse. Figure 1.10 shows the different forest types and their locations within Australia, highlighting that much of ~~the-our~~<sup>mh</sup> forested lands are frequently found near population centres along the east coast. 16% of Australia is covered by forest, most (75%) of which is Eucalyptus, a major emitter of VOCs (Guenther et al. 2012).

It has been estimated by [the MEGAN biogenic emission model \(Section 1.3.2\)](#) [MEGAN<sup>tb</sup>](#) that Australian vegetation is among the world's strongest isoprene emitters, with forests in SE Australia having emission factors greater than  $16 \text{ mg m}^{-2} \text{ h}^{-1}$  (see figure 1.11) (Guenther et al. 2006; Guenther et al. 2012). Measurement campaigns in SE Australia have since cast doubt on the emission factors used by MEGAN, potentially due to poor characterisation of Eucalyptus trees and soil moisture (Emmerson et al. 2016; Emmerson et al. 2019). These emissions factor estimates are not well verified and measurements of isoprene (or other BVOC) emissions are sparse and infrequent in Australia (Sindelarova et al. 2014; Bauwens et al. 2016).

Australia has a much greater diversity of tree species than is represented by MEGAN, however sparse measurements of emissions and BVOC concentrations makes model improvement difficult. Uncertainties in isoprene emissions could explain why models of HCHO over Australia are poor at reproducing satellite measurements (Stavrakou et al. 2009).

Australia suffers from poor characterisation of plant emissions, partly because many of the emission factors are based on northern hemispheric data. Many plant emissions rates have not been published, such as those for any Australian acacias. Some Eucalypt emissions are based on samples from young trees, which may emit more isoprene than older trees (Emmerson et al. 2016). Changes in parameterisation of soil moisture in the MEGAN model lead to large reductions (38 – 58%) in Australian isoprene emission estimates, although errors remain (Sindelarova et al. 2014; Emmerson et al. 2019).

The only VOC emission information currently available come from three campaigns: the Measurements of Urban, Marine and Biogenic Air (MUMBA), and the two Sydney Particulate Studies (SPS1 and SPS2) (see Section 2.2). These measurements take place in south eastern Australia, and have suggested lower isoprene emissions than seen in models (by a factor of 2-6) (Emmerson et al. 2016). Several improvements are required as no simple scaling factor can completely fix the misrepresentation of isoprene emissions (Emmerson et al. 2016). These measurements focus on air quality and biogenic VOCs and use several different instruments (including PTR-MS and GC-FID) to detect metrics such as air particulates, HCHO, isoprene, and meteorological information. There is also an instrument at University of Wollongong (see Section 2.2.3.3) that provides 20 years of HCHO measurements, which is the only available long term HCHO vertical column measurement record in Australia comparable to satellite data. Satellite HCHO column measurements can be limited by various factors including interfering species, water, clouds, orography, etc, and independent in situ measurements are required to validate the data. For further details on these campaigns and measurements, see Section 2.2. [Improvements to emissions models require improved understanding of emissions from Australian vegetation and how they respond to meteorological and chemical parameters. However, constraining](#)

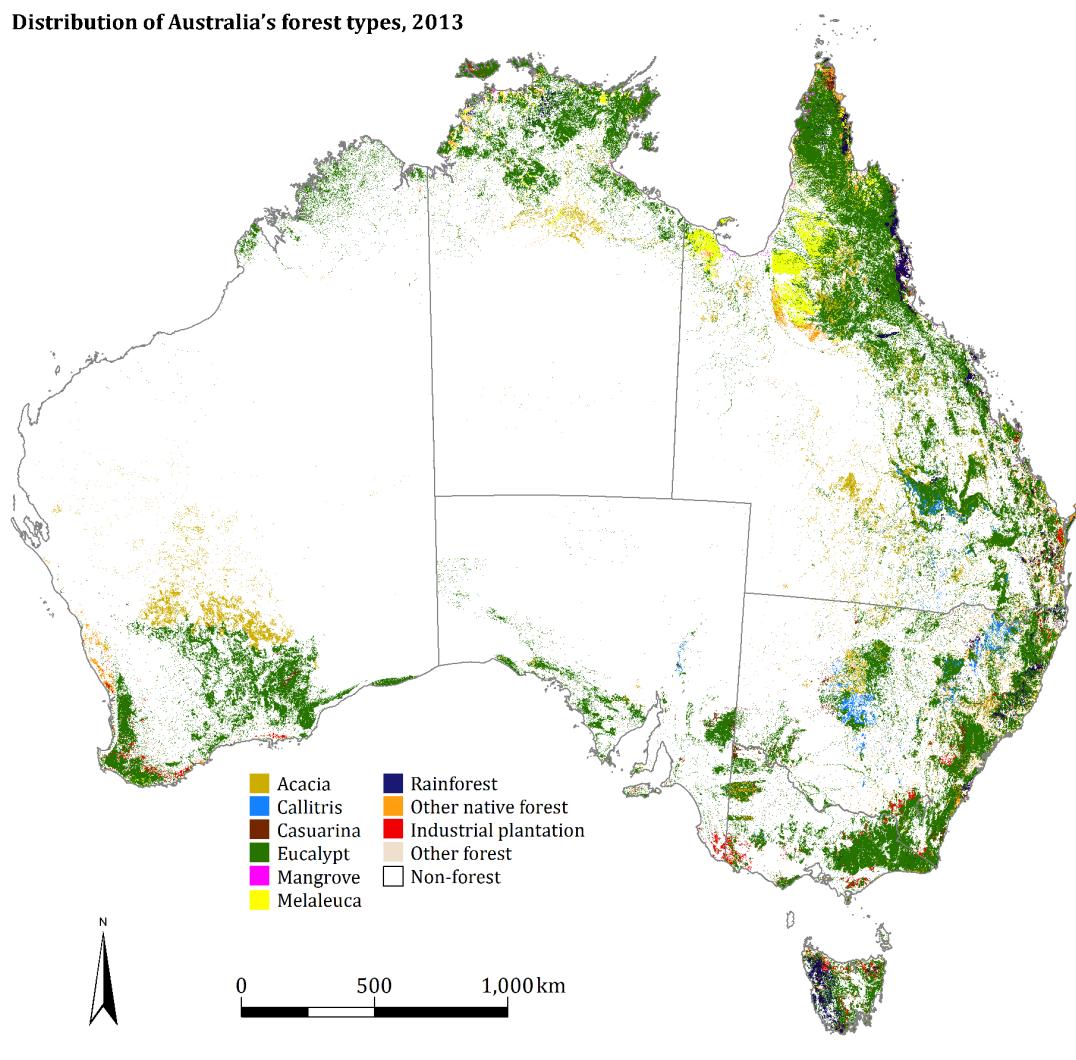


FIGURE 1.10: Forest types in Australia reproduced from <http://www.agriculture.gov.au/abares/forestsaustralia/australias-forests>.

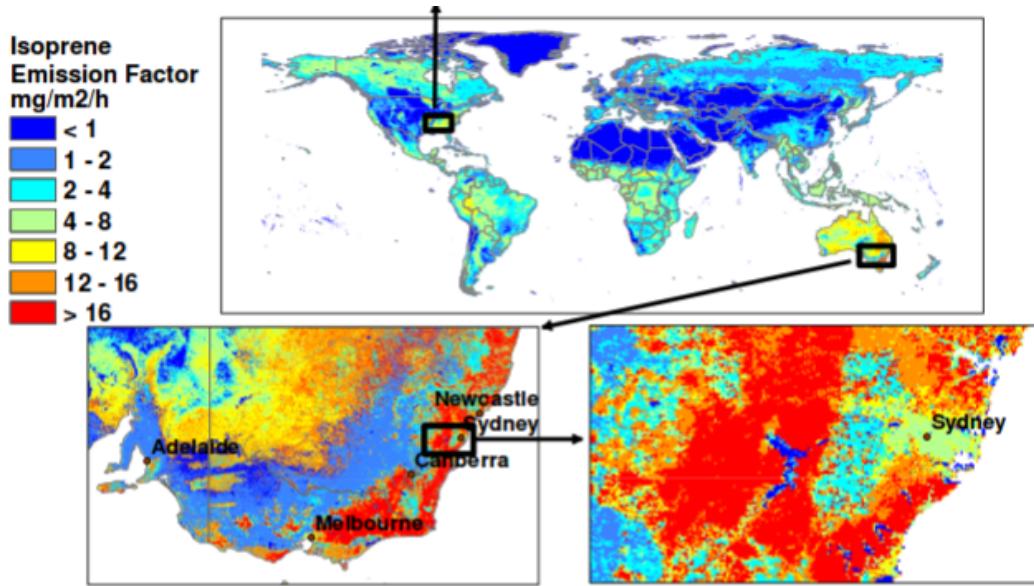


FIGURE 1.11: Global and Australian isoprene emission factors modified from Guenther et al. (2006).

these processes is difficult given the lack of available isoprene measurements.<sup>mh</sup> Satellite measurements of HCHO can instead be used to estimate and improve Australian isoprene emissions without costly measurement campaigns. This is due to the near-linear relationship between HCHO and isoprene emissions (e.g., Palmer et al. 2001; Millet et al. 2006; Bauwens et al. 2016). This method is explored in this thesis as described in Chapter 3.

## 1.7 Summary of gaps in knowledge

Several layers of knowledge are required to accurately estimate ozone concentration in the troposphere, with the most important being emissions of precursors, transport from the ozone rich stratosphere, and chemical processes in both clean and polluted air masses. These natural processes that lead to ozone production are poorly understood in Australia mostly due to the difficulty and expense involved in measuring both ozone and precursors over any representative sample of Australia. Improvements to emissions models require improved understanding of emissions from Australian vegetation and how they respond to meteorological and chemical parameters. How Australian tropospheric ozone concentrations are affected by ozone transported from the stratosphere is also unclear, largely due to a lack of knowledge around how frequently a transport event occurs, and the quantity of ozone transported by any such event. Finally, due to the complex nature of atmospheric chemistry involving isoprene and other BVOCs, it is not currently known how sensitive ozone concentrations in the atmosphere are to changes in BVOC emissions.<sup>mh</sup>

## 1.8 Aims

**This thesis aims In this thesis I aim<sup>mh</sup> to improve understanding of natural contributions to ozone over Australia and the southern ocean.** The two largest contributors to tropospheric ozone concentrations are chemical production (driven by precursor emissions) and transport from the stratosphere. **This thesis aims I aim<sup>mh</sup> to improve understanding of both of these sources using existing satellite and ground-based datasets along with modelled outputs from the GEOS-Chem chemical transport model.**

Estimation of BVOC emissions in Australia can be improved through satellite measurements of HCHO, one of isoprene's primary oxidation products. Satellites that overpass daily record measurements over all of Australia. Combining satellite data with model outputs provides a platform for the understanding of natural processes, **which are uncertain over Australia<sup>mh</sup>**. Satellite measurements require modelled a priori vertical profiles of HCHO to estimate total column amounts. **This thesis aims I aim<sup>mh</sup> to recalculate satellite vertical columns of HCHO using updated model a priori information.** In this effort **I aim to improve the understanding of<sup>mh</sup>** the importance of relevant parameters (within GEOS-Chem) in calculating vertical columns of HCHO measured by satellite **is analysed<sup>mh</sup>**. This includes an examination of how well GEOS-Chem simulates several species such as NO<sub>x</sub>, isoprene, and HCHO compared to both in situ and remote measurement data that exists for Australia. Additionally, **I detail<sup>mh</sup> the construction and effects of satellite data filters is examined<sup>mh</sup>**. The work towards this aim is detailed in Chapter 2.

The technique of determining isoprene emissions from satellite detected HCHO is called satellite inversion. **This thesis aims I aim<sup>mh</sup> to determine isoprene emissions in Australia using a top-down inversion of satellite HCHO, through a modelled yield from isoprene to HCHO.** HCHO amounts and the yield of isoprene to HCHO over Australia is required to create top-down estimates. This process also requires careful examination of when the assumptions required within the inversion process are not valid. Due to the low availability of in situ data over most of the Australian continent, a combination of modelled and satellite data can reduce the uncertainties of isoprene emissions from Australian landscapes. Improved emissions estimates will in turn improve the accuracy of CTMs, providing better predictions of atmospheric composition and its response to ongoing environmental change. The work towards fulfilling this aim is in Chapter 3.

Stratospheric transport is the second largest driver of tropospheric ozone concentrations, and an improved understanding of transported ozone can be determined from ozonesonde measurements. **This thesis aims I aim<sup>mh</sup> to improve understanding of ozone transported to the troposphere from the stratosphere in Australia and the southern ocean.** Ozonesondes provide a glimpse of the vertical ozone profile up to  $\sim 30$  km. **In this thesis, and I use<sup>mh</sup> a Fourier filter is used<sup>mh</sup> to determine how often stratospheric transport is occurring at three sites: Melbourne, Macquarie Island, and Davis Station.** Transport event frequency analysis combined with modelled ozone distributions is used to derive a new method of detection and quantification of transported ozone as described in Chapter 4.

Overall, **this thesis aims I aim<sup>mh</sup> to describe relative importance of sources of**

tropospheric ozone in Australia, as well as their seasonality. This includes a description of I-will-describe<sup>mh</sup> how modelled ozone is affected by updated isoprene emissions, a comparison of comparing<sup>mh</sup> changes in GEOS-Chem outputs, and an analysis of by<sup>mh</sup> ozone transport from the stratosphere.<sup>mh</sup> This thesis will provide new insight into tropospheric ozone in Australia.

## Chapter 2

# Data and Modelling

### 2.1 Introduction

In this thesis the word model is used in lieu of chemical transport model (CTM), a class of model that simulates chemistry and chemical transport through the atmosphere. Models of the atmosphere can be used to interpret measurements, estimate chemical concentrations at any scale, and predict atmospheric composition in the future. A more detailed discussion of CTMs can be found in Section 1.5.2.<sup>tb</sup> Some advanced measurement techniques make use of modelled a priori information in order to produce useful outputs. Models of ozone in the atmosphere are used broadly for international assessments of ozone precursor emissions, and estimating effects from related processes (such as radiation) (Young et al. 2017). Models provide an estimate of many trace gas concentrations; however, verification is required, and is generally performed using results from measurement campaigns. In situ measurements from campaigns or measurement stations can be used to examine what is happening at a particular location. These data are used to determine how accurate models or estimates are – however the utility is limited to where and when the measurements took place.<sup>tb</sup> In this thesis data from several campaigns are compared against model outputs and satellite datasets. Satellite datasets provide large amounts of data over most of the planet. However, they can have high uncertainty due to instrument limitations. In this chapter several satellite datasets are combined to estimate biogenic HCHO amounts over Australia.

The first aim is to analyse Australia-specific HCHO amounts measured by satellite. These are then used along with model data to estimate isoprene emissions, this is the second aim and takes place in Chapter 3. The third aim is to quantify ozone transported from the stratosphere down into the troposphere (Chapter 4). The focus in this chapter is to describe and analyse model outputs and measurements along with how they are calculated and compared for use in achieving the aims of this thesis. Section 2.2 details satellite and campaign datasets and describes model outputs. Measurement techniques used to retrieve the most utilised satellite dataset are outlined in Section 2.4. Section 2.3 describes the GEOS-Chem model, how it is run and what setup and outputs are used in this thesis. In Section 2.5 the process of using model outputs to recalculate satellite vertical columns is described and analysed. In order to compare satellite data with other datasets, some work must be undertaken to avoid introducing bias (e.g., Palmer et al. 2001; Eskes and Boersma 2003; Marais et al. 2012; Lamsal et al. 2014). One key step is to recalculate the satellite information using modelled data, detailed in Section 2.6. The effects of these recalculations on satellite HCHO is also

examined. [The creation and effects of filters used to remove non-biogenic influences are described in Section 2.7.](#)<sup>tb</sup>

## 2.2 Datasets

This section describes the datasets used in this thesis, along with an overview of the measurement techniques used for each. This includes modelled output, satellite measurements, and measurement campaigns. These datasets serve four purposes:

1. Model output validation is performed in this chapter
2. Calculation of biogenic HCHO distribution over Australia is performed in this chapter
3. Recalculated OMI formaldehyde columns are used as a basis for estimating isoprene emissions in Chapter 3
4. Model outputs and measurements are used quantify ozone transport in Chapter 4

Details on filtering and interpolations which are undertaken when reading data are also provided, as each dataset has its own resolution. While no measurements are made throughout this thesis, it is important to understand the techniques used to create utilised datasets in order to understand possible anomalous data points or trends.

Horizontal geographical coordinates are always discussed in terms of degrees north and east, from -180° to 180° spanning west to east around the globe and -90° to 90° spanning the latitudes from the Antarctic to the Arctic. Vertical resolution is sometimes discussed in terms of metres, sometimes in terms of pressure (hPa), and sometimes in terms of sigma ( $\sigma$ ) coordinates. Sigma coordinates represent the fraction of the atmosphere vertically above, with  $\sigma = 1$  being the surface and  $\sigma = 0$  being the top of the atmosphere (TOA). This can be useful when running global atmospheric models as the ground altitude is always at  $\sigma = 1$  and [topography is not an issue we need not worry about topography](#)<sup>mh</sup>. Conversion between  $\sigma$  and pressure (p) coordinates is as follows:

$$\sigma = \frac{p_s - p}{p_s - p_t} \quad (2.1)$$

where  $p_s$  and  $p_t$  is surface pressure and pressure at the TOA respectively.

Uncertainty (or error) is present in each dataset and where possible the causes are explained. There are two types of error: systematic and random. Arguably the worst of these is systematic error (or bias). Bias normally indicates a problem in calculation or instrumentation. If the systematic error is known, it can be corrected for by either offsetting data in the opposite direction, or else fixing the cause. A proper fix can only be performed if the sources of error are known and there is a way of correcting or bypassing them. Random error is often reported as some function of a datasets variance, or uncertainty. It can be reduced through averaging either spatially or temporally. Temporal and/or spatial averaging decreases uncertainty by a factor of  $1/\sqrt{N}$  where N is the number of observations being averaged.

### 2.2.1 Satellite

Satellite data products are generally classed into several categories, level zero through to level three. level zero products are sensor counts and orbital swath data, level one-B data calibrates and geo-locates the level zero data. level two products additionally have temporal, spatial, solar, and viewing geometry information, as well as quality flags. level three data is a temporally and spatially aggregated subset of level two data, for instance monthly or yearly averages.

Difficulties can arise when aerosols (e.g., clouds, smoke, dust) interfere with recorded spectra; however, some of these can be detected and filtered out. Instruments including MODIS (onboard the Aqua and Terra satellites) are able to determine aerosol optical depth (AOD), a measure of atmospheric scatter and absorbance. An AOD under 0.05 indicates a clear sky, while values of 1 or greater indicate increasingly hazy conditions. This is important in order to determine where measurements from other instruments may be compromised by high interference. Cloud filtering is performed on several satellite products used in this thesis, which reduces uncertainty at the cost of measurement quantity. This has been seen to introduce a clear-sky bias in monthly averages since measurements do not include cloudy days (Surl, Palmer, and Abad 2018).

Satellite instruments measuring atmospheric composition of interest in this thesis generally record spectra between around 250-700 nm split into spectral components at around 0.3 nm (e.g., Leue et al. 2001). This allows calculation of the abundance of trace gases including O<sub>3</sub>, NO<sub>2</sub>, and HCHO. Satellite measurements are generally performed using spectral fitting followed by conversion to vertical column densities. Several public data servers are available which include products from satellites, including NASA's Earthdata portal (<https://earthdata.nasa.gov/>) and the Belgian Institute for Space Aeronomy (IASB-BIRA) Aeronomie site (<http://h2co.aeronomie.be/>). The following subsections describe some of the satellite products used in this thesis.

#### 2.2.1.1 Formaldehyde

Satellite formaldehyde data are used in this thesis to constrain Australian isoprene emissions (Chapter 3).<sup>mh</sup> HCHO products can be found in four satellite instruments: GOME (Global Ozone Monitoring Experiment) on ERS-2 (second European Remote Sensing satellite), SCIAMACHY (SCanning Imaging Absorption SpectroMeter for Atmospheric CHartographY) on ENVI-SAT (ENVIronmental SATellite), OMI (Ozone Monitoring Instrument) on the Aura satellite, and GOME2 on MetOp-A and MetOp-B (Meteorological Operational satellites). These satellites have slightly different spectral and spatial resolutions, as well as using varied processes to estimate HCHO from detected radiances. This leads to different measurements between instruments (Lorente et al. 2017), and both validation and comparison become more important when using these remotely sensed data. The dataset used in this thesis is from the OMI onboard the Aura satellite, as it has data over the time period covered by GEOS5 modelled meteorological information used in GEOS-Chem (Section 2.3), and sufficiently covers the southern hemisphere.

Satellite based formaldehyde measurements from the OMI instrument onboard Aura are stored in the OMHCHO product. OMHCHO data is used and modified

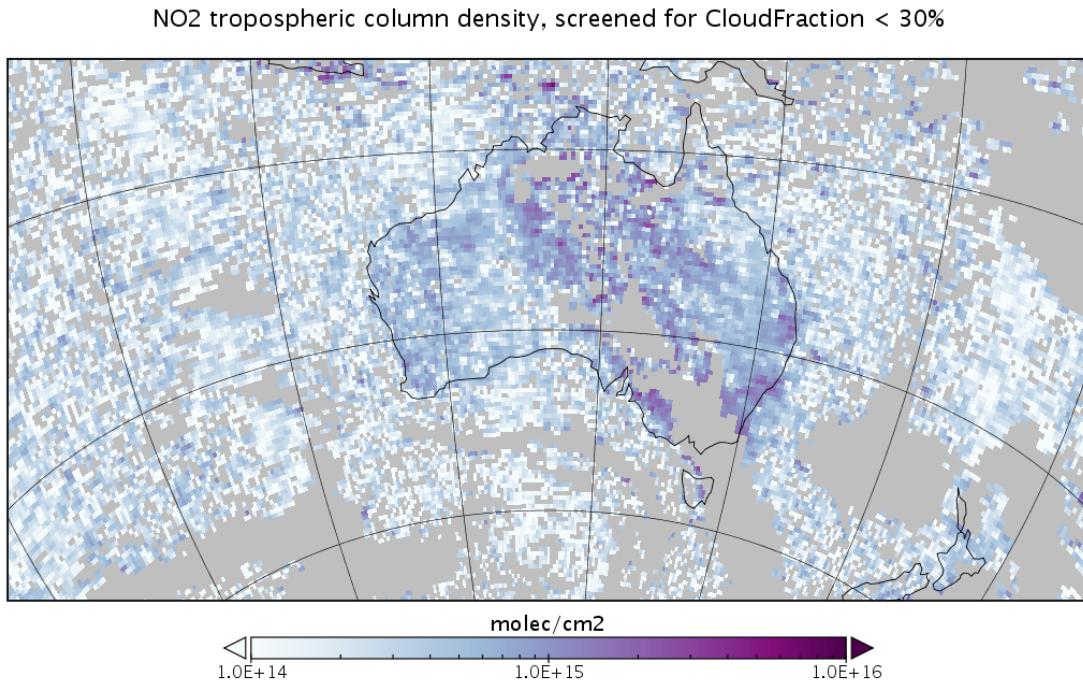


FIGURE 2.1: Example of NO<sub>2</sub> tropospheric columns for 29 Jan. 2005 from the OMNO2d product. Data downloaded from <https://search.earthdata.nasa.gov/search>, DOI:10.5067/Aura/OMI/DATA3007.

extensively throughout this thesis, and so is discussed in more detail in Section 2.4. Calculation of column density and AMF are discussed respectively in sections 2.4 and 2.4.2.

### 2.2.1.2 Nitrogen dioxide

Daily OMI NO<sub>2</sub> (OMNO2d) is a gridded daily level three NO<sub>2</sub> product of satellite measurements averaged into  $0.25^\circ \times 0.25^\circ$  horizontally resolved bins. An example from Jan 29, 2005 is shown in Figure 2.1, while an average for 2005 (global) is shown in Figure 2.2. The measurement resolution is 40 km by 130 km. NO<sub>2</sub> measured by OMI is used in this thesis to check whether NO<sub>2</sub> is well represented by GEOS-Chem (see Section 2.3.5 for the comparison between this product and GEOS-Chem calculations). It is also used to form the anthropogenic influence filter for calculating biogenic-only HCHO columns (see Section 3.2.3.2).

Like other satellite products, OMNO2d is influenced by a priori model profiles, which are required to convert slant path radiance to vertical columns. These models are generally low resolution ( $\sim 110 \text{ km} \times 110 \text{ km}$ ), which leads to column smearing and difficulty detecting point sources of high NO emissions (Goldberg et al. 2017). Uncertainty in this product arises mostly from the calculation of the AMF (up to 50% of total error) (Lorente et al. 2017).

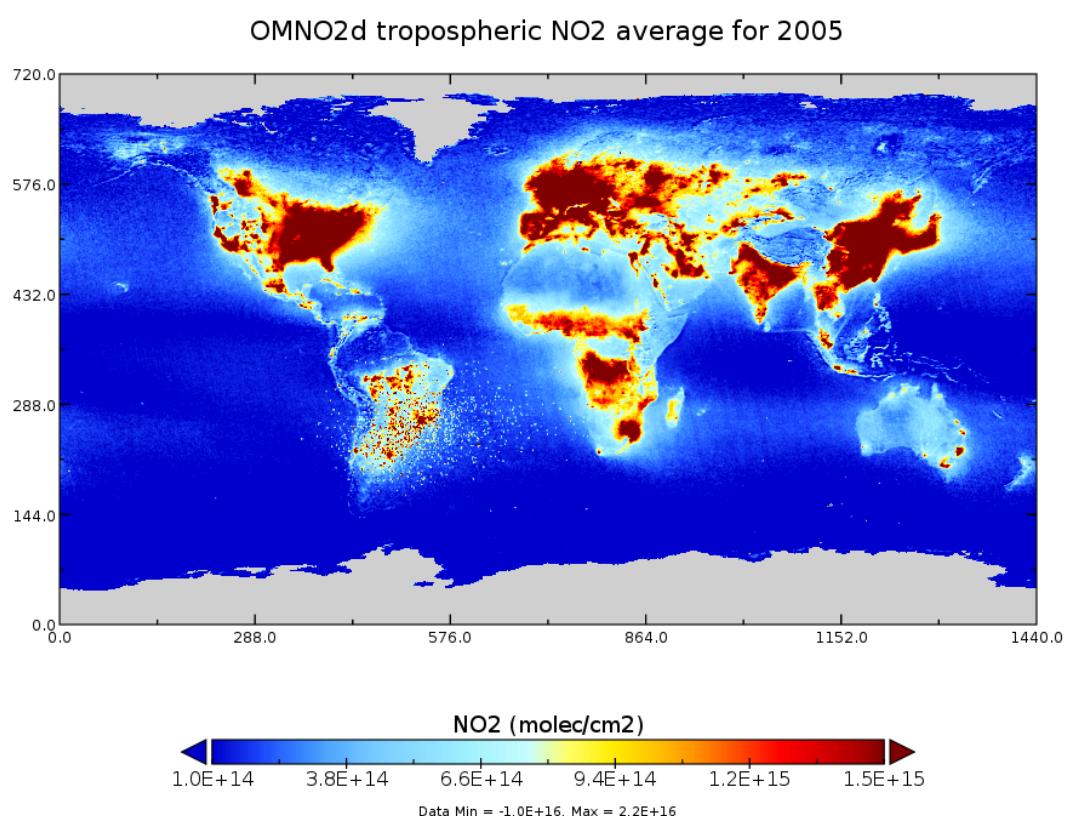


FIGURE 2.2: Average 2005 tropospheric NO<sub>2</sub> from OMNO2d with pixels screened for < 30% cloud cover.

### 2.2.1.3 Aerosol optical depth

Aerosols in the atmosphere can be seen through their interactions with visible radiation. Smoke and dust can be seen as an increase in aerosol optical depth (AOD) (see Section 1.4.2.2). This is due to these particles scattering and absorbing UV radiation (Ahn, Torres, and Bhartia 2008). A data product provided by Earthdata ([https://disc.gsfc.nasa.gov/datasets/OMAERUVd\\_V003/summary](https://disc.gsfc.nasa.gov/datasets/OMAERUVd_V003/summary)) called OMAERUVd (DOI: 10.5067/Aura/OMI/DATA3003) is used in this thesis to filter out satellite HCHO data that may be influenced by pyrogenic emissions (see Section 3.2.3.1)<sup>mh</sup>.

OMAERUVd allows detection of areas which may be smoke-affected. The product contains AOD and aerosol absorption optical depths (AAOD) at three wavelengths (354, 388, and 500 nm), along with UV aerosol index. The OMAERUVd product is level three, based on quality filtered level two swath pixels which are then gridded by daily averaging. These products are most sensitive to error in the form of sub-pixel scale cloud interference, however AAOD is the less sensitive of the two (Ahn, Torres, and Bhartia 2008). In this thesis AAOD is used to filter out potential smoke-affected areas.

In this work AAOD is mapped from  $1^\circ \times 1^\circ$  horizontal resolution to  $0.25^\circ \times 0.3125^\circ$  using nearest value mapping. The AAOD at 500 nm wavelength is used to determine smoke influence, although any of the provided wavelengths would be affected by smoke plumes and could also be used. Any areas with daily AAOD > 0.03 are considered to be potentially smoke plume affected (see Section 3.2.3.1).

### 2.2.1.4 Active fires

Filtering pyrogenic influence from satellite HCHO requires knowledge of where fire is occurring, which is determined using the MOD14A1 dataset as applied in Section 3.2.3.1.<sup>mh</sup> MOD14A1 is a gridded daily satellite dataset of fire counts at  $1 \times 1 \text{ km}^2$  horizontal resolution. Fire counts are observed four times daily from Terra (10:30, 22:30 local time) and Aqua (01:30, 13:30 local time). The fire pixels are detected based on parameters including apparent pixel temperature and the nearby background temperature. The dataset is obtained from NASA Earth Observations that is part of the EOS Project Science Office at the NASA Goddard Space Flight Center [https://neo.sci.gsfc.nasa.gov/view.php?datasetId=MOD14A1\\_M\\_FIRE](https://neo.sci.gsfc.nasa.gov/view.php?datasetId=MOD14A1_M_FIRE). This product is downloaded and binned into the lower  $0.25^\circ \times 0.3125^\circ$  resolution (using the sum of fire pixels) to create an active fire influence mask (see Section 3.2.3.1)<sup>mh</sup>.

### 2.2.1.5 Carbon monoxide

In Chapter 4, potential biomass burning plumes are identified using satellite observations of carbon monoxide (CO) from the AIRS (Atmospheric Infra-red Sounder) instrument aboard the Aqua satellite (Texeira 2013). The Aqua satellite overpasses at approximately 01:30 and 13:30 local time, with NASA producing level three gridded output at  $1^\circ \times 1^\circ$ . CO is used as a proxy for biomass burning plumes, and used to qualitatively attribute ozone intrusion events (Section 4.3.5).

### 2.2.1.6 Uncertainties

While satellite data is effective at covering vast areas (the entire earth) it only provides information at a particular time of day, is affected by cloud cover, and generally does not have fine vertical resolution. Measurements retrieved by satellites have large uncertainties, which arise from the process of transforming spectra into total column measurements, as well as from instrument degradation (satellite instruments are hard to repair once they are launched). Uncertainty in transforming satellite spectra comes from a range of issues, including measurement difficulties introduced by clouds, and instrument sensitivity to particular aerosols (Millet et al. 2006). Many products require analysis of cloud and aerosol properties in order to estimate total column amounts (Palmer et al. 2001; Palmer et al. 2003; Marais et al. 2012; Vasilkov et al. 2017).

## 2.2.2 Model datasets

### 2.2.2.1 GEOS-Chem output

The GEOS-Chem model is used extensively in this thesis and is discussed in more detail in Section 2.3. GEOS-Chem model outputs are described in Section 2.3.6.1. These are generally resolved to 47 vertical levels from the ground up to 0.01 hPa, at  $2^\circ \times 2.5^\circ$  horizontal resolution.

### 2.2.2.2 Meteorological reanalysis

Synoptic scale weather patterns are taken from the European Centre for Medium-range Weather Forecasts (ECMWF) Interim Reanalysis (ERA-I) (Dee et al. 2011). These are used in Chapter 4 to determine typical weather systems for stratospheric ozone intrusions. The ERA-I output was the most up to date at the time (2016) but has since been superseded by ERA5.

### 2.2.2.3 Surface temperatures

The Climate Prediction Center (CPC) provides a product with maximum daily land-surface temperature at  $0.5^\circ \times 0.5^\circ$  horizontal resolution. These data are used to check the correlation between HCHO and temperature at a higher resolution than is provided by GEOS-Chem output. A full description of the data can be found at <https://www.esrl.noaa.gov/psd/data/gridded/data.cpc.globaltemp.html>. CPC Global Temperature data is provided by National Oceanic and Atmospheric Administration/Office of Oceanic and Atmospheric Research/Earth System Research Laboratory Physical Sciences Division, Boulder, Colorado, USA, from their web site at <https://www.esrl.noaa.gov/psd/>. An example of one day of land temperature output is shown in Figure 2.3.

## 2.2.3 In situ and ground based datasets

Data from several in situ measurement campaigns and long-term measurement programs are used to examine the accuracy of modelled data at specific sites. Figure 2.4

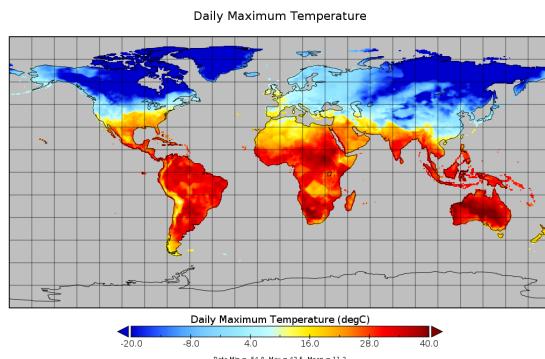


FIGURE 2.3: Climate Prediction Center daily maximum temperature data for 1 Jan. 2005.

TABLE 2.1: Detection limits for MUMBA

Dates	HCHO (ppb)	Isoprene (ppb)	Ozone (ppb)
21/Dec/2012 - 29/Dec/2012	0.205	0.003	0.5
29/Dec/2012 - 18/Jan/2013	0.105	0.005	0.5
19/Jan/2013 - 15/Feb/2013	0.186	0.003	0.5

shows the locations of ground-based measurement sites in the top panel, and release sites for ozonesondes in the bottom panel.

### 2.2.3.1 Short term measurement campaigns

Some campaign datasets (described in this section) provide separate time series for brief periods of both isoprene and formaldehyde. Figure 2.5 shows these along with the detection limits and also shows isoprene measurements superimposed over a single year. It is apparent that more measurements are required to see more than the daily cycles.

The MUMBA campaign (Paton-Walsh et al. 2017) measured various compound abundances including isoprene, formaldehyde, and ozone from 21 December 2012 to 15 February 2013. These measurements took place in Wollongong, 10 m above ground level (40 m above sea level). Ozone was measured by Thermo UV absorption with 1-minute time resolution averaged into hourly outputs. Isoprene and HCHO were measured by Ionicon Proton-Transfer-Reaction Mass spectrometer (PTR-MS), with a time resolution of 3-minutes, averaged each hour. Detection limits varied due to instrument conditions, and are listed in Table 2.1. The full dataset has been published on PANGAEA (DOI:10.1594/PANGAEA.871982) (Guérette et al. 2018).

Isoprene concentration measurement uncertainty in this product is estimated by Dunne et al. (2018) to be 50%. Although the uncertainty determined through calibration measurements was only 15% (Guérette et al. 2018), this does not account for competing trace gas interference (such as furan). When used in this thesis, readings are re-sampled to hourly averages, and measurements below the detection limit are set to half of the detection limit. The major sources of uncertainty include interference

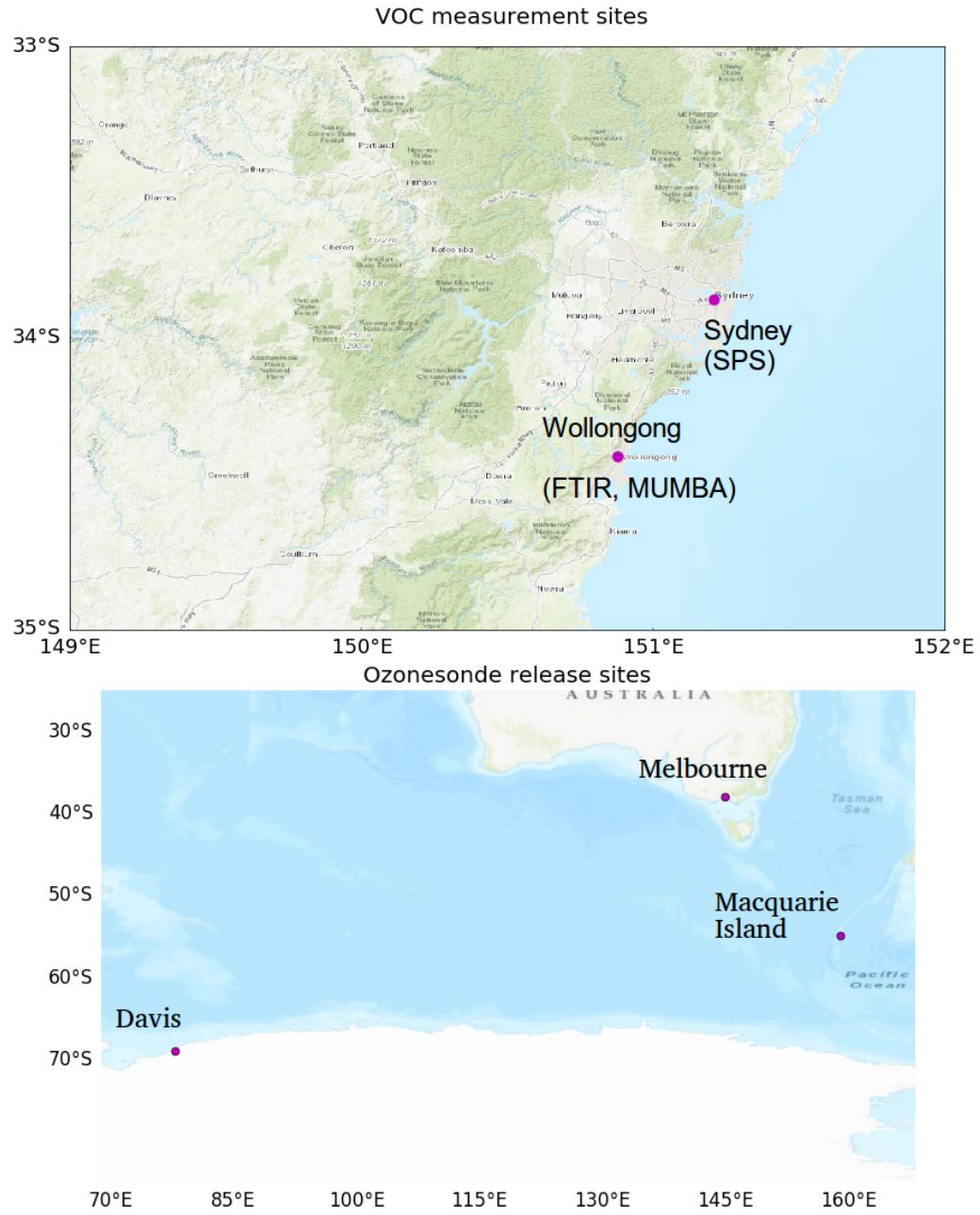


FIGURE 2.4: Locations of ground based in situ measurements (top panel) and ozonesonde release sites (bottom panel).

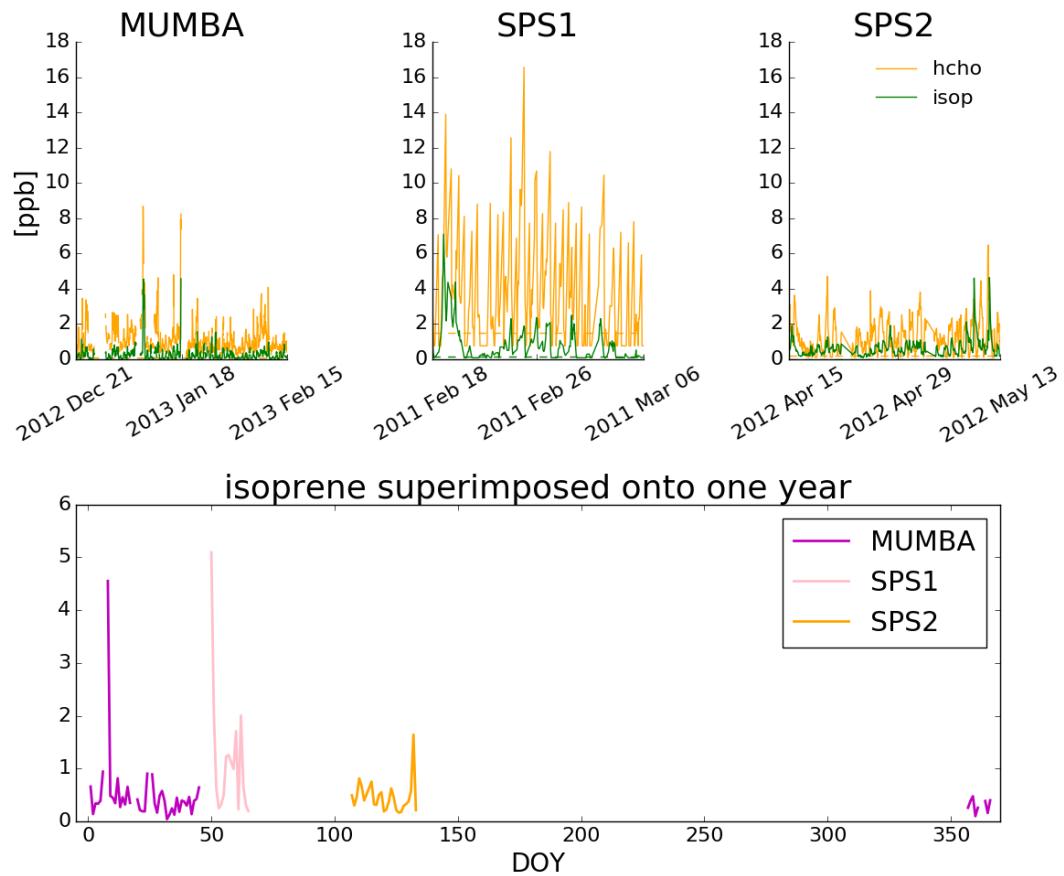


FIGURE 2.5: Top: MUMBA, SPS1, and SPS2 time-series for HCHO (orange) and isoprene (green), along with detection limits (dashed). Bottom: isoprene measurements superimposed onto a single year.

from non-target compounds and under-reporting (e.g., Dunne et al. 2018; Guérette et al. 2018).

Two trace gas measurement campaigns took place at the Westmead air quality station: stage 1 (SPS1) from 5 February to 7 March, 2011 and stage 2 (SPS2) from 16 April to 14 May, 2012. Two instruments measured VOC concentrations: a PTR-MS, and a gas chromatograph with a flame ionisation detector (GC-FID). The PTR-MS uses chemical ionisation mass spectrometry and can quantify VOCs at high temporal resolution (< 1 s). It was calibrated several times per day against HCHO, isoprene,  $\alpha$ -pinene, and several other VOCs. Further measurement specifics can be found in Dunne et al. (2018).

The output provides hourly averaged concentrations of trace gases based on the mass to charge ratio ( $m/z$ ), which for isoprene is 69. It is possible that other chemicals (such as furan, with the same  $m/z$ ) interfered with this value, especially at low ambient isoprene concentrations and towards the end of autumn (SPS2) when wood fires start to become frequent (Guérette et al. 2018). The GC-FID analysed samples collected in multi-absorbent tubes, with lower temporal resolution but no interference. Further details for this method can be found in Cheng et al. (2016). GC-FID data are averaged from 05:00-10:00 local time, and 11:00-19:00 local time, while PTR-MS data are averaged hourly. Significant differences occur between measurement devices when detecting isoprene, potentially due to interfering compounds in the PTR-MS (Dunne et al. 2018).

### 2.2.3.2 Ozonesondes

Ozonesonde data come from the World Ozone and Ultraviolet Data Centre (WOUDC). Ozonesondes are weather balloons that measure ozone concentrations using an electrochemical concentration cell (<http://www.ndsc.ncep.noaa.gov/organize/protocols/appendix5/>). Ozonesondes provide a high vertical resolution profile of ozone, temperature, pressure, and humidity. Generally the instrument will perform 150-300 measurements from the surface to around 35 km. Ozonesondes are launched approximately weekly from Melbourne (38° S, 145° E), Macquarie Island (55° S, 159° E) and Davis (69° S, 78° E). More information on this dataset is given in Section 4.3.

### 2.2.3.3 Wollongong FTIR

Upon the roof of the Chemistry building at the University of Wollongong lies a solar Fourier transform infra-red spectrometer (FTIR) which measures HCHO (amongst other gases) in the path of light between the instrument and the sun. The instrument is part of the Network for the Detection of Atmospheric Composition Change (NDACC) and data can be retrieved from the NDACC database (<http://www.ndaccdemo.org/stations/wollongong-australia>). The current principle investigator producing and quality assuring the dataset is Dr. Nicholas Jones.

Measurements of vertical profiles are affected by the instrument's sensitivity to HCHO at all vertical levels. Vertical mixing ratios resolved to 48 vertical levels are retrieved. The averaging kernel ( $A$ ) essentially represents the sensitivity of the retrieval technique to HCHO concentrations at different altitudes. Figure 2.6 shows averaged retrievals  $x_{ret}$  and the a priori side by side. Most of the HCHO is close to the surface,

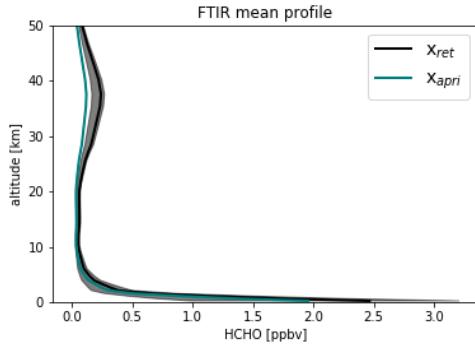


FIGURE 2.6: Mean midday FTIR profile ( $x_{ret}$ ) and a priori profile ( $x_{apri}$ ). Shaded area shows the inter-quartile range over the same time period. Profile is averaged between November 2007 and April 2013.

and generally the a priori lies within the inter-quartile range of the retrievals, just below the mean. Figure 2.7 shows how the retrievals of total vertical column HCHO ( $\Omega$ ) are sensitive to HCHO between 0 km and 60 km altitude. The panel on the right shows how each retrieved vertical level is sensitive to HCHO at all levels. This shows there is not much sensitivity at higher altitudes and HCHO at lower altitudes can affect retrieved values for higher altitudes. When comparing FTIR measurements to modelled data, vertical columns are used rather than profiles as the degrees of freedom in each measurement is low ( $\sim 1.3$  in summer to  $\sim 1.7$  in winter). Degrees of freedom signify how many pieces of information can be retrieved from a measurement, and profiles resolved to  $N$  vertical levels would require at least  $N-1$  degrees of freedom.

The retrieved vertical profile ( $x_{ret}$ ) can be expressed as a linear function of the true vertical profile ( $x_{true}$ ) and the a priori vertical profile ( $x_{apri}$ ):

$$x_{ret} = x_{apri} + A(x_{true} - x_{apri}) \quad (2.2)$$

One way to understand Equation 2.2 is to consider that if the device was perfectly sensitive to HCHO concentrations at all vertical levels (making  $A = I$  the identity matrix), then the retrieval would be equal to the true profile. If the instrument was perfectly insensitive (making  $A = 0$ ), then the retrieval would completely ignore the true profile and simply equal the a priori. To compare modelled vertical profiles ( $x_{GC}$ ) with the FTIR, the modelled profiles are *convolved* with the instrument averaging kernel and a priori to create a pseudo retrieval ( $x'_{GC}$ ) which shows what the instrument would retrieve if the model were the truth:

$$x'_{GC} = x_a + A(x_{GC} - x_a) \quad (2.3)$$

$x'_{GC}$  can be compared against  $x_{ret}$  to determine if measurements and modelled vertical profiles or columns show significant differences. This convolution is required so as to avoid introducing a bias between the instrument and the model which is due to the instrument lack of sensitivity. Retrieved vertical profiles can be transformed into vertical columns ( $\Omega$ ) using the hydrostatic relationship and an assumed dry air profile

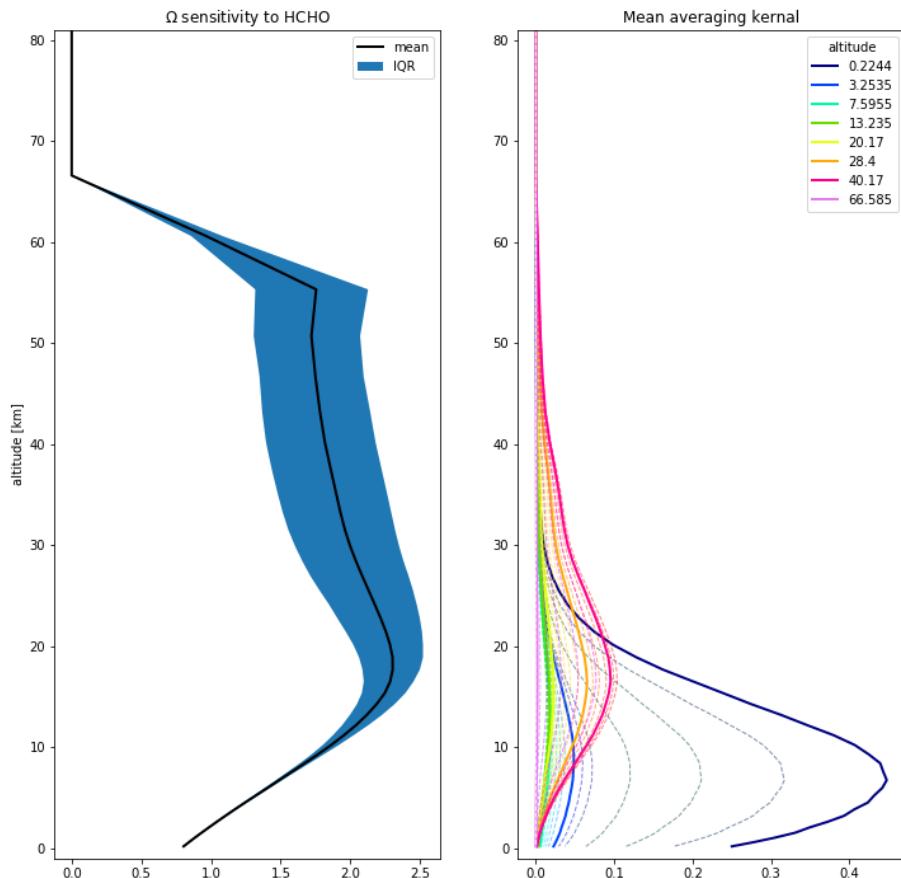


FIGURE 2.7: Left panel shows the mean midday (13:00 - 14:00 local time) total column averaging kernel, along with the inter-quartile range (IQR) between November 2007 and April 2013. Right panel shows the mean averaging kernel for the vertical profile over the same time period, coloured by vertical level. One in six vertical levels are labelled and plotted with a solid line, and the rest are shown with dashed lines.

The x-axes are unitless in the two plots shown here.

(see Deeter (2002)):

$$\Omega = x \times 2.12 \times 10^{13} \times \Delta p \quad (2.4)$$

where  $\Omega$  is in molec cm<sup>-2</sup>,  $x$  is in ppbv, and  $\Delta p$  is the pressure in hPa between pressure midpoints for each vertical level in  $x$ .

## 2.3 GEOS-Chem

### 2.3.1 Overview

GEOS-Chem is a global, Eulerian CTM (see Section 1.5.2) with a state of the science chemical mechanism, and transport driven by meteorological input from the Goddard Earth Observing System (GEOS) of the NASA Global Modelling and Assimilation Office (GMAO). Chemistry, transport, and meteorology are simulated at 15 minute time steps within a global set of 3-D boxes. Emissions are either prescribed by inventories (e.g., fire emissions are prescribed from the global fire emissions database GFED4) or modelled (e.g., biogenic emissions are modelled using the Model of Emissions of Gases and Aerosols from Nature (MEGAN)).

GEOS-Chem simulates more than 100 chemical species within the atmosphere, from the earth's surface up to 0.01 hPa. Modelled concentrations can be used in combination with measurements to give a verifiable estimate of atmospheric gases and aerosols in places and times where no measurements exist. It was developed, and is maintained, by Harvard University staff as well as users and researchers worldwide. In this thesis [GEOS-Chem version 10.01 is used](#)~~I use version 10.01 of GEOS-Chem<sup>mh</sup>~~, which outputs up to 66 chemical species (tracers) in the standard run, at  $2^\circ \times 2.5^\circ$  horizontal resolution, with 47 levels up to the top of the atmosphere (TOA at 0.01 hPa).

Global CTMs are often run using one or several emission models (or the output from them) to determine input fluxes. Some of the underlying model data used by GEOS-Chem are described here. Meteorological parameters are taken from NASA's GEOS-5 dataset ( $0.5^\circ \times 0.666^\circ$ ) (Chen et al. 2009), which exists from December 2003 to April 2013. GEOS-5 meteorological fields are used to drive atmospheric transport. Fire emissions come from the global fire emissions database (GFED4) product (Giglio, Randerson, and Van Der Werf 2013). Anthropogenic VOC emissions come from the Emission Database for Global Atmospheric Research (EDGAR) inventory, while biogenic VOC emissions are simulated using the MEGAN model (see Section 2.3.4).

### 2.3.2 Installing and running GEOS-Chem

GEOS-Chem instructions for download, compilation, and running can be found in the user guide provided by Harvard: <http://acmg.seas.harvard.edu/geos/doc/man/>. In order to build and run GEOS-Chem a high-speed computing system is optimal, as globally gridded chemical calculations can take a long time to perform ([for us](#)~~mh~~~70 computation hours per month [in this work<sup>mh</sup>](#)). For this thesis, GEOS-Chem is [installed](#)~~I installed~~ GEOS-Chem<sup>mh</sup> onto a suitably configured workspace on the Raijin supercomputer at the National Computational Infrastructure (NCI, <http://nci.org.au/>). This workspace included access to compilers and libraries that are needed to

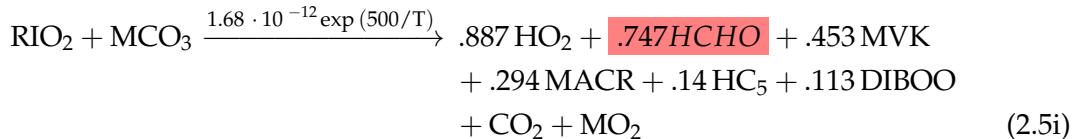
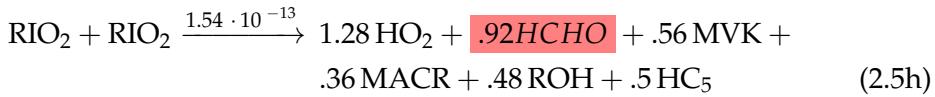
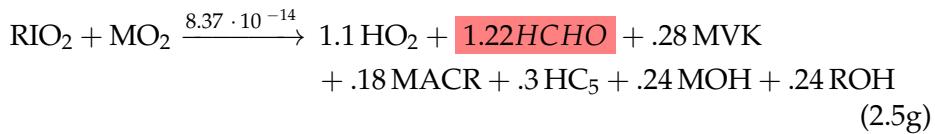
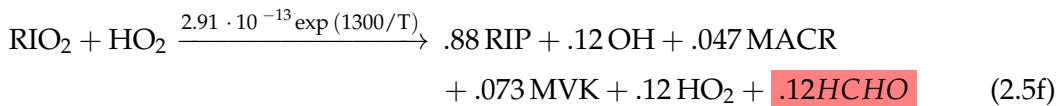
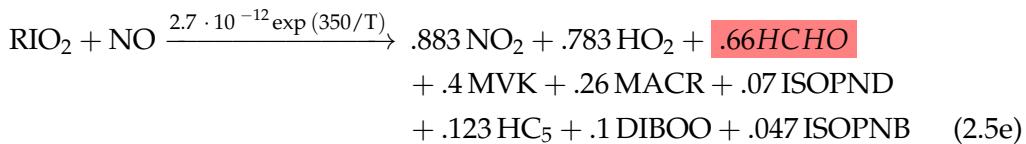
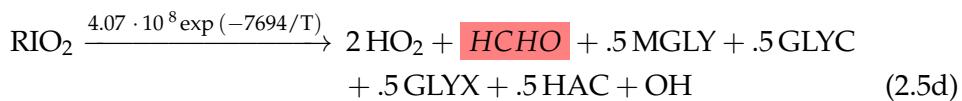
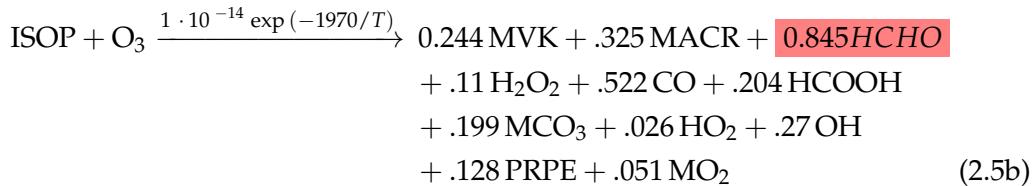
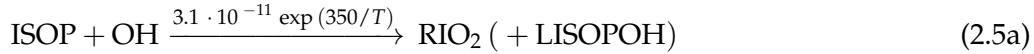
build the Fortran based GEOS-Chem source code, and IDL, Python, and various editors and scripting languages to read, run, edit, and analyse both GEOS-Chem and its output.

### 2.3.3 Chemical Mechanism

A chemical mechanism is a closed system of chemical reactions and their associated rate constants. Chemical reactions are represented by systems of differential equations to be solved for each gridbox in GEOS-Chem. Simplifications are required due to the large number of reactions that occur in the atmosphere, and the coupled and stiff nature of these reactions that slow down computation of the solutions (Brasseur and Jacob 2017). Mathematical stiffness in chemical systems of differential equations is due to the massively differing reaction time scales—for instance hydroxyl radicals react within seconds while methane has an atmospheric lifetime of 8–10 years (Wuebbles and Hayhoe 2002).<sup>b</sup>

This section focuses on isoprene oxidative chemistry simulated by GEOS-Chem, as it will be used to determine the yield from isoprene emissions to HCHO columns.

Some of the important reactions involving isoprene are reproduced here, including reaction rates ( $k$ ) in the form  $k = A \exp(-E/RT)$ , where  $T$  is temperature,  $E$  is activation energy, and  $R$  is the gas constant.  $A$ ,  $E$ , and  $R$  are predefined for each reaction. The following set of equations (2.5) lists the main isoprene and related reactions, with terms defined in Table 2.2. LISPOH is added in order to allow the model to keep track of how much isoprene is oxidised by OH. The reactions shown here are just the first-stage oxidations of isoprene and the isoprene peroxy radical from OH oxidation, subsequent reactions will also produce HCHO. The mechanism used in version 10.01 (and its history) is described online at [http://wiki.seas.harvard.edu/geos-chem/index.php/NOx-Ox-HC-Aer-Br\\_chemistry\\_mechanism](http://wiki.seas.harvard.edu/geos-chem/index.php/NOx-Ox-HC-Aer-Br_chemistry_mechanism).



The isoprene reactions simulated by GEOS-Chem were originally described by Horowitz et al. (1998). The mechanism was subsequently updated by Mao et al. (2013), who changed the isoprene nitrate yields and added products based on Paulot et al. (2009a) and Paulot et al. (2009b). They used the yields and reactions of various positional isomers of isoprene nitrates, and their oxidation products. Further mechanistic properties, like isomerisation rates, are based on results from four publications: Peeters, Nguyen, and Vereecken (2009), Peeters and Muller (2010), Crounse et al. (2011), and Crounse et al. (2012).

In part, GEOS-Chem chemistry mechanisms are based on atmospheric chemistry chamber studies. In a chamber with clean air and high NO concentrations, isoprene photooxidation is initially driven by OH addition, followed by NO<sub>x</sub> chemistry (150 min – 600 min), and finally HO<sub>x</sub> dominated chemistry. Formation of isoprene nitrates

TABLE 2.2: Species and lumped species from the GEOS-Chem mechanism.

Name	Definition
ACTA	Acetic acid: CH <sub>3</sub> C(O)OH
DIBOO	Dibble peroxy radical
GLYC	Glycoaldehyde: HOCH <sub>2</sub> CHO
GLYX	Glyoxal: CHOCHO
HAC	Hydroxyacetone: HOCH <sub>2</sub> C(O)CH <sub>3</sub>
INO2	RO <sub>2</sub> from ISOP+NO <sub>3</sub>
ISNP	an isoprene nitrate
ISOPNB	$\beta$ isoprene nitrates
ISOPND	$\delta$ isoprene nitrates
MACR	Methacrolein: CH <sub>2</sub> =C(CH <sub>3</sub> )CHO
MCO3	Peroxyacetyl radical: CH <sub>3</sub> C(O)OO
MEK	Methyl ethyl ketone: RC(O)R
MGLY	Methylglyoxal: CH <sub>3</sub> COCHO
MO2	Methylperoxy radical: CH <sub>3</sub> O <sub>2</sub>
MOH	Methanol: CH <sub>3</sub> OH
MVK	Methylvinylketone: CH <sub>2</sub> =CHC(=O)CH <sub>3</sub>
PRPE	>C <sub>3</sub> alkenes: C <sub>3</sub> H <sub>6</sub> , ...
RIO2	isoprene peroxy radical: ROO
ROH	>C <sub>2</sub> alcohols

(ISOPN) affect ozone levels through NO<sub>x</sub> sequestration, the yields and fate of these nitrates was analysed in Paulot et al. (2009a). Prior to 2012, oxidation chamber studies were performed in high NO or HO<sub>2</sub> concentrations, giving peroxy lifetimes of less than 0.1 s (Crounse et al. 2012; Wolfe et al. 2012). In most environments NO and HO<sub>2</sub> concentrations are not so high.<sup>tb</sup> GEOS-Chem uses production rates for different NO concentrations and peroxy radical lifetimes determined by Mao et al. (2013). Crounse et al. (2011) examined the isomerisations associated with the oxidation of isoprene to six different isomers of ROO formed in the presence of oxygen.<sup>tb</sup> The primary oxidation pathway of isoprene is reaction with the OH radical (ISOP + OH → RIO<sub>2</sub>) shown in Equation 2.5a. Isoprene undergoes OH addition at the 1 and 4 positions, becoming  $\beta$  (71%) or  $\delta$  (29%) ROO, although these are not distinguished in the v10 GEOS-Chem mechanism except in the distribution of the isoprene nitrates. Additional oxidative pathways are from ozonolysis (Equation 2.5b) and reaction with NO<sub>3</sub> (Equation 2.5c). These pathways are much slower but can be important in particular scenarios such as inside a pollution plume or at night when NO<sub>3</sub> radicals can build up. Following isoprene oxidation, seven potential ROO reactions (RIO<sub>2</sub> in Equation 2.5) can occur depending on reactant concentrations and local temperature. Reactions 2.5d - 2.5j compete to determine the fate of ROO, which is often grouped into high or low NO<sub>x</sub> concentration pathways.

### 2.3.3.1 Nitrogen oxide impacts

Isoprene reactions that occur following emission GEOS-Chem reactions do not distinguish between high and low NO<sub>x</sub> scenarios. However, when discussing the likely reactions isoprene undergoes after emission, the dominant reactions generally depend on NO<sub>x</sub> levels. This leads to isoprene reactions being broadly discussed within the context of high or low NO<sub>x</sub> environmental conditions. High NO<sub>x</sub> ( $\sim 1$  ppb) has been defined as conditions where NO reactions are the main cause of losses for ROO (Reaction 2.5e) (Palmer et al. 2003).

In low NO<sub>x</sub> environments, ROO losses occur from isomerisation or reaction with several compounds. NO<sub>x</sub> + HO<sub>2</sub> (Reaction 2.5f) produces mostly hydroxy hydroperoxides (ISOPOOH), and some HCHO. ISOPOOH can be oxidised (by OH) to produce epoxydiols, recycling OH (Paulot et al. 2009b). Isomerisation of ROO (Reaction 2.5d) largely produces HCHO, while recycling OH and producing HO<sub>2</sub>. This isomerisation accounts for 1,5-H shifts producing MACR, MVK, HCHO, or 1,6-H shifts producing HPALDs. HPALDs can photolysis to regenerate OH and small VOCs (Crounse et al. 2011; Wolfe et al. 2012; Jozef et al. 2014). In low NO<sub>x</sub> environments, ROO oxidation has a lower HCHO yield as reaction with HO<sub>2</sub> yields 12% HCHO (compared to 66% when reacting with NO) (Mao et al. 2013). Refer to Section 1.3.4 for more information. Less frequently, ROO reacts with itself, MO<sub>2</sub>, or MCO<sub>3</sub>.

Under high NO<sub>x</sub> conditions, the fate of ROO differs depending on how it was formed. The  $\beta$ -hydroxyl reacts with NO<sub>x</sub> and produces HCHO (66%), methylvinylketone (40%) (MVK), methacrolein (26%), and  $\beta$ -hydroxyl nitrates (6.7%) (ISOPNB). The  $\delta$ -hydroxyl reacts with NO to form  $\delta$ -hydroxyl nitrates (24%) (ISOPND), and ISOPNB (6.7%). These two pathways are combined in Reaction 2.5e. The numbers in this section are determined in Mao et al. (2013) and form the basis for some of GEOS-Chem chemical mechanism.

### 2.3.3.2 OH regeneration

Isoprene chemistry in GEOS-Chem includes OH regeneration from oxidation of epoxydiols (not shown in Equation 2.5) and slow isomerisation of ROO (Equation 2.5d) (Mao et al. 2013). This regeneration dealt with the problem seen in older models where ISOPOOH production titrated OH, which was not backed up by measurements (Paulot et al. 2009b; Mao et al. 2013). Until recently, models struggled to represent OH in the atmosphere, due to missing chemistry and issues in measurement techniques. Prior to Mao et al. (2012), measurements of OH in high VOC regions may have been up to double the real atmospheric OH levels, due to formation of OH inside measuring instruments.<sup>tb</sup> OH regeneration through photolysis of HPALDs in areas with high isoprene emissions are included in GEOS-Chem based on Peeters and Muller (2010). Photolysis of photolabile peroxy-acid-aldehydes generates OH, improving model agreement with continental observations. OH and HPALD interactions are central to maintaining the OH levels in pristine and moderately polluted environments, which makes isoprene both a source and sink of OH (Peeters and Muller 2010; Taraborrelli et al. 2012). Mao et al. (2013) showed that drastically lowering (by a factor of 50) the rate constant for ROO isomerisation lead to better organic nitrate agreements with measurements. These chemical updates led to more accurate modelling of OH concentrations, especially in low NO<sub>x</sub> conditions common in remote forests.<sup>tb</sup> The updates to isoprene chemistry

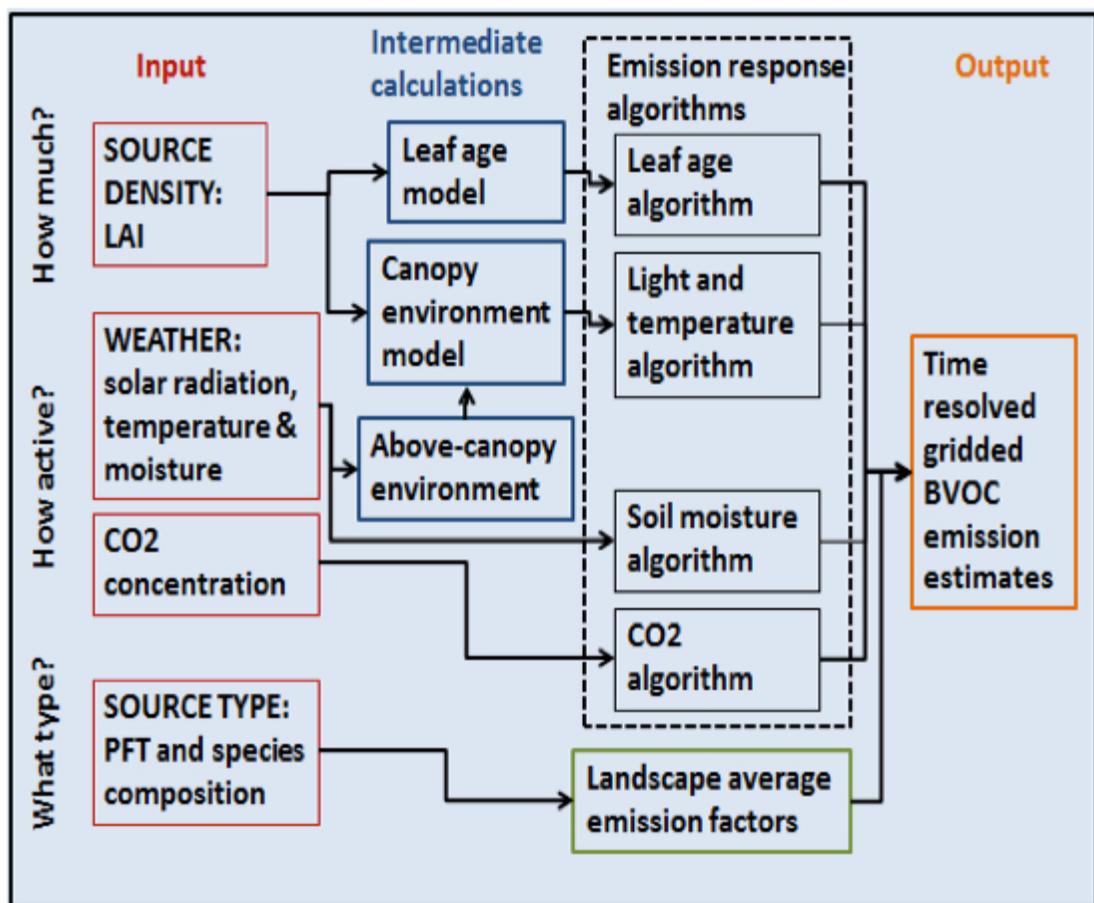


FIGURE 2.8: MEGAN schematic, from <http://lar.wsu.edu/megan/>. In GEOS-Chem, a pre-computed emission factors map is used for isoprene in lieu of the soil moisture algorithm.

by Mao et al. (2013), and those shown in Crounse et al. (2011) and Crounse et al. (2012) are the last implemented in the version of GEOS-Chem used in this thesis.

### 2.3.4 Biogenic emissions from MEGAN

GEOS-Chem includes online coupling to the Model of Emissions of Gases and Aerosols from Nature (MEGAN) to determine biogenic emissions globally. GEOS-Chem V10.01 uses MEGAN V2.1 described by Guenther et al. (2012). MEGAN uses leaf area index (LAI), global meteorological data, and photosynthetic photon flux density to simulate terrestrial isoprene emissions (Kefauver, Filella, and Peñuelas 2014). In GEOS-Chem MEGAN uses high resolution emission factor maps described in Guenther et al. (2012). The schematic for MEGAN, from <http://lar.wsu.edu/megan/>, is shown in Figure 2.8.

MEGAN was developed as a replacement for two earlier canopy-environment emission models (the Biogenic Emission Inventory System, and the Global Emissions Initiative), initially including a simple canopy radiative transfer model that parameterised sun-lit and shaded conditions through a canopy. Early models did not account for abiotic stresses, such as drought, prior rainfall and land use changes. These stresses

influence species-specific emissions by more than an order of magnitude (Niinemets et al. 1999). Isoprene emissions were based on temperature, leaf area, and light, but have since been updated to include leaf age activity (Guenther et al. 2000), and a leaf energy balance model (Guenther et al. 2006) in MEGANv2.0. This update included a parameter for soil moisture, to account for drought conditions; however, this parameter is currently (as of version 2.1) not applied in GEOS-Chem (Sindelarova et al. 2014). Instructions to run version 2.1 are available at [http://lar.wsu.edu/megan/docs/MEGAN2.1\\_User\\_GuideWSU.pdf](http://lar.wsu.edu/megan/docs/MEGAN2.1_User_GuideWSU.pdf). Version 2.1 (updated from 2.0 (Guenther et al. 2006)) includes 147 species, in 19 BVOC classes, which are lumped for mechanisms in chemical models. GEOS-Chem uses an embedded version of MEGAN 2.1.

GEOS-Chem computes some emissions using predefined emission factor ( $\epsilon$ ) maps from MEGAN source code, and others using PFT maps and associated emission factor relationships. For isoprene (the focus in this thesis) MEGAN calculates emissions online, using local meteorological conditions. Emissions  $E$  of species  $i$  are calculated using these EF for classes of plant types  $j$  and associated grid-box coverage  $\chi$  and an activity factor  $\gamma$ , which accounts for response to environmental conditions. The following equation is reproduced from Guenther et al. (2012) showing how emissions  $E_i$  are determined in MEGAN:

$$E_i = \gamma_i \sum_j \epsilon_{i,j} \chi_j \quad (2.6)$$

For example: isoprene emissions are tied to radiation, temperature, plant emission strength ( $\epsilon$ ), and leaf area index, for each grid box. In GEOS-Chem the isoprene emission factors (EF) are pre-calculated and the equation becomes  $E_{\text{isoprene}} = \gamma_{\text{isoprene}} EF$ .

### 2.3.5 Nitrogen oxides

$\text{NO}_x$  concentrations affect atmospheric oxidative capacity, which changes many factors important in estimating isoprene emissions including isoprene to HCHO yield, isoprene lifetime, and isoprene oxidation pathways. In GEOS-Chem,  $\text{NO}_x$  concentrations are regulated by  $\text{O}_3$ , VOC,  $\text{HO}_x$ , bromine and aerosols reactions.  $\text{NO}_x$  emissions from agriculture, power generation, and combustion based transport drive enhancements in populated areas. In GEOS-Chem, anthropogenic emissions are taken from the Emissions Database for Global Atmospheric Research (EDGAR, for more details visit [http://wiki.seas.harvard.edu/geos-chem/index.php/EDGAR\\_v4.2\\_anthropogenic\\_emissions](http://wiki.seas.harvard.edu/geos-chem/index.php/EDGAR_v4.2_anthropogenic_emissions)). Other  $\text{NO}_x$  emissions arise from sources including soil emissions and lightning. Soil emissions are laid out in Hudman et al. (2012), parameterised using biome specific emission factors, and an explicit fertiliser dataset. Lightning based  $\text{NO}_x$  production is created based on inventories of lightning flash rates scaled within GEOS-Chem. Conversion to nitric acid ( $\text{HNO}_3$ ) followed by deposition is the primary  $\text{NO}_x$  removal mechanism (Delmas, Serca, and Jambert 1997; Ayers and Simpson 2006).

In order to determine the accuracy of GEOS-Chem simulated  $\text{NO}_x$  over Australia, modelled  $\text{NO}_2$  amounts are compared to satellite data (where available) for 2005. Figure 2.9 shows tropospheric  $\text{NO}_2$  columns from GEOS-Chem, OMI satellite data OMNO2d, and their differences over four seasons in 2005. Simulated GEOS-Chem tropospheric  $\text{NO}_2$  columns averaged from 13:00-14:00 local time are compared against

OMNO2d data that are averaged into seasonal  $2^\circ \times 2.5^\circ$  bins. Both datasets show Sydney and Melbourne as NO<sub>2</sub> hotspots; however, Sydney itself is underestimated by GEOS-Chem throughout the year, likely due to low or poorly aligned emission maps (shown in Figure 2.10). Over much of the country GEOS-Chem overestimates NO<sub>2</sub> by 10-60%, except in the northern areas where up to 50% underestimation occurs in summer. An underestimation in southern NSW can also be seen in the later half of the year. Notably GEOS-Chem anthropogenic NO does not have any seasonality (Figure 2.10), while soil shows clear differences in emission hotspots between summer and winter.

Figure 2.11 shows scatter plots with one data point for each land square over Australia, for each season, coloured by total NO emissions. The correlation between model and satellite NO<sub>2</sub> columns is reasonable throughout the year over Australia, with a general positive bias throughout the year in modelled amounts. A comparison between the bias (GEOS-Chem - OMNO2d) with anthropogenic or soil emissions (columns 2 and 3 respectively) shows no real correlation, suggesting the bias is not driven by either anthropogenic or soil NO emissions in any season. Without a clear link between emissions and biases, alterations would be too subjective, and a change to NO<sub>x</sub> chemistry is beyond the scope of this thesis. Additionally, high satellite NO<sub>2</sub> columns are filtered in later calculations that assume biogenic air masses as these high NO<sub>2</sub> columns suggest anthropogenic influence. The conclusion drawn is that modelled anthropogenic and soil NO emissions do not show sufficient evidence of biasing GEOS-Chem NO<sub>2</sub> columns away from satellite measurements over Australia. For this reason modelled NO emissions are not modified in model runs in this thesis.

### 2.3.6 GEOS-Chem simulations

GEOS-Chem is run five times independently in this thesis, with different outputs from each simulation used to determine specific information. The different output types are first described in Section 2.3.6.1. Following this is the list of model runs, including a summary of the run, outputs created, and a summary of how they are used (Section 2.3.6.2). Finally a brief comparison between a subset of the runs is performed (Section 2.3.6.3).

#### 2.3.6.1 GEOS-Chem outputs

GEOS-Chem in this thesis is run with a 15 minute time step for transport and 30 minutes for chemistry, at  $2^\circ \times 2.5^\circ$  horizontal resolution over 47 vertical levels. Output is the average over many of these time steps. Optionally one to many columns can be output at high temporal resolution. This feature has been used here to compare modelled ozone with ozonesonde profiles at three sonde release sites discussed in Chapter 4.

**Midday output** is output from averaging over a window of local time for each grid-box. Output averaged between 13:00-14:00 local time is saved to match with Aura satellite measurements, as Aura overpasses at  $\sim$ 1330 local time each day. Midday output is saved both for comparison with, and recalculation of, satellite measurements.

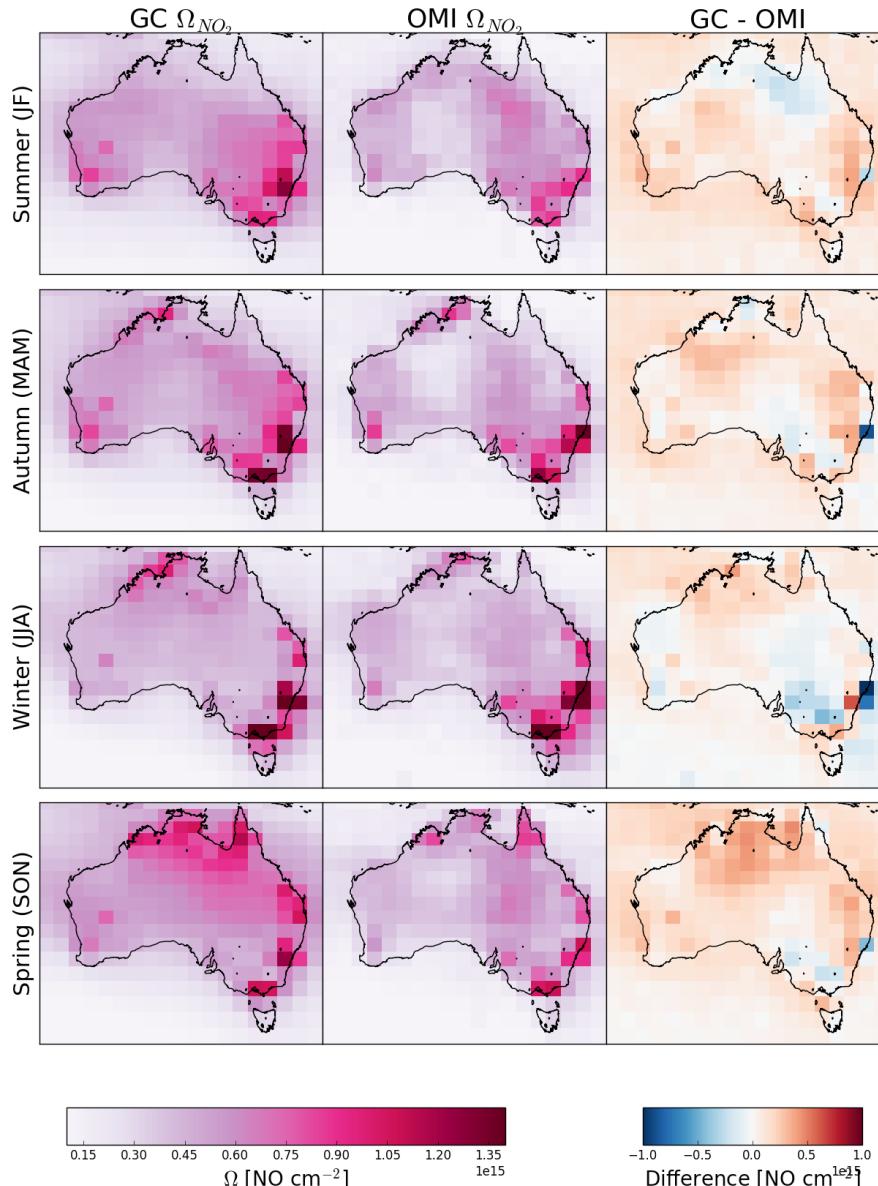


FIGURE 2.9: Left to right columns show tropospheric NO<sub>2</sub> columns ( $\Omega_{NO_2}$ ; molec  $cm^{-2}$ ) from GEOS-Chem (GC), OMNO2d (OMI), and the differences. Each row shows one season from 2005, the left two columns use the left colour scale, while the third column uses the right colour scale.

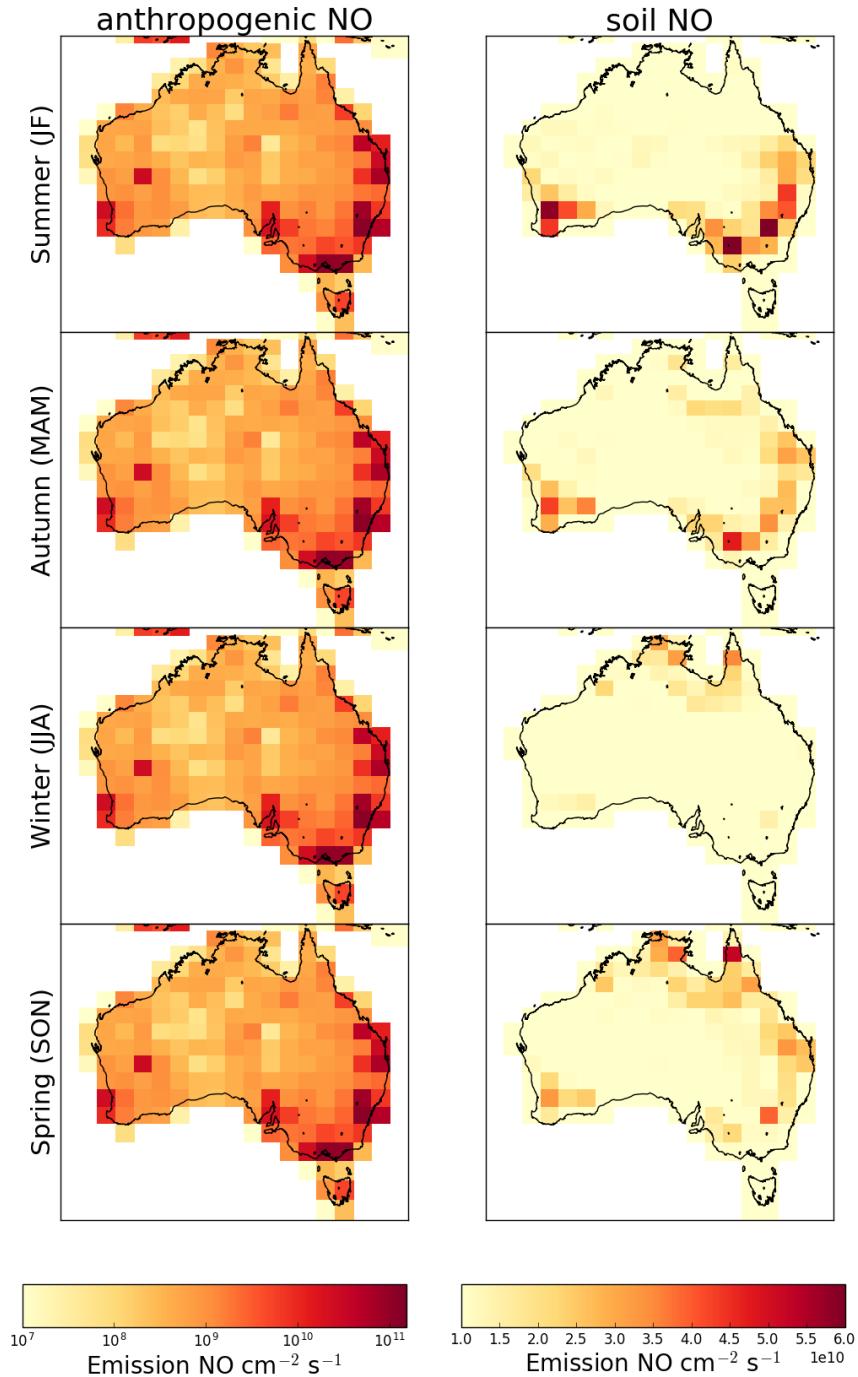


FIGURE 2.10: Left and right columns show anthropogenic and soil emissions of NO respectively from GEOS-Chem in molec  $\text{cm}^{-2} \text{s}^{-1}$ . Each row shows one season from 2005. Anthropogenic and soil emissions use a logarithmic and linear colour scale respectively.

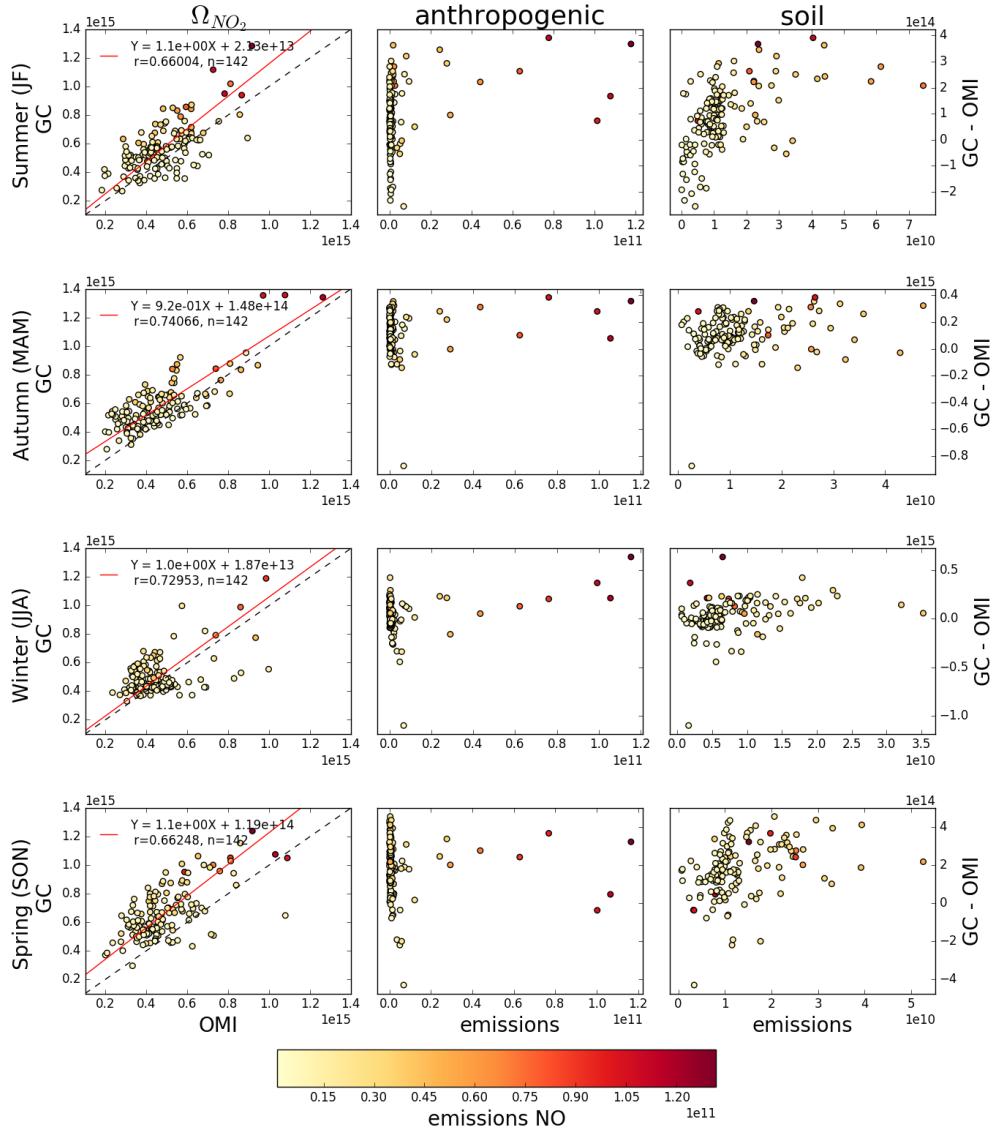


FIGURE 2.11: The first column shows scatter plots of tropospheric  $\text{NO}_2$  columns between GEOS-Chem (y-axis) and OMNO2d (x-axis) at  $2^\circ \times 2.5^\circ$ . The reduced major axis linear regression is drawn in red, and the equation, regression coefficient ( $r$ ), and number of grid squares ( $n$ ) used is inlaid as a legend. The second and third columns show the scatter between emissions and the bias between GEOS-Chem and OMNO2d (GC-OMI), for anthropogenic and soil emissions respectively. The third column has a different x-axis scale for each season. These right two columns share the far right axis; however, emissions are from anthropogenic and soil sources respectively. All scatter points are coloured by the sum of anthropogenic and soil NO emissions (from GEOS-chem), as per the colour bar shown at the bottom.

**HEMCO diagnostics** the Harvard-NASA Emissions Component (HEMCO) deals with emissions inventories used in GEOS-Chem. When working with globally gridded data, handling local time offsets becomes more important. The hourly averaged emissions of isoprene are saved using GMT, and an offset based on the longitude is used to retrieve values at local time. This offset is determined as one hour per  $15^\circ$  (as  $360^\circ$  is 24 hours).

**Tracer averages** are daily or monthly averaged gridbox concentrations.

**Time series** are vertical profiles of ozone and other species and meteorological diagnostics saved at a temporal resolution of up to 15 minutes.

One issue with modelled outputs over Australia is that they are difficult to compare to in situ measurements as they are averaged over a large horizontal space and vertical volume.

### 2.3.6.2 GEOS-Chem runs

The following list summarises each model run as well as enumerating the outputs (described above), and how the run is used in the thesis.

**UCX** The model was run using the Universal tropospheric-stratospheric Chemistry eXtension (UCX) mechanism with 72 vertical levels from the surface to the top of the atmosphere (TOA  $\sim 0.1$  hPa). UCX runs a chemistry mechanism with combined tropospheric and stratospheric reactions, with an increased number of stratospheric calculations performed online relative to the default (tropospheric chemistry) simulation (Eastham, Weisenstein, and Barrett 2014).

1. Outputs: Midday output, daily tracer averages, and time series over three stations.
2. This run shows what influence the stratospheric chemistry additions have over tropospheric isoprene, HCHO, and ozone concentrations.
3. The daily tracer averages are used to determine ozone intrusion quantification (Section 4.6.1), and ozone concentration seasonality (Section 4.5).
4. Time series outputs are compared against ozonesonde releases (Section 4.5) both over time and vertically.

**Tropchem (standard)** default settings for GEOS-Chem 10.01, using 47 vertical levels at  $2^\circ \times 2.5^\circ$  horizontal resolution. Additional midday output is created to allow AMF recalculation code to run on OMI satellite measurements (Section 2.6.3.2).

1. Outputs: Midday output, daily tracer averages, and HEMCO diagnostics
2. These are used in recalculation of the satellite AMF (Section 2.5), and the modelled background HCHO over the remote Pacific that is used in the reference sector correction for OMI column retrievals (Section 2.6.5). Additional diagnostic outputs are added by the author to allow AMF recalculation.

3. Midday output is combined from two different runs in order to determine smearing of the isoprene-to-HCHO relationship (Section 3.2.3.3)
4. Total yearly ozone concentrations are compared before and after scaling isoprene emissions using the top-down estimate.

**Tropchem (isoprene emissions halved)** identical to standard tropchem except isoprene emissions are halved.

1. Outputs: Midday output, and monthly tracer averages
2. They are used to check modelled ozone sensitivity to isoprene emissions (following section Figure 2.16).
3. They are also combined with the standard run to determine model sensitivity to transport (Section 3.2.3.3)

**Tropchem (biogenic emissions only)** identical to standard tropchem except all non-biogenic emissions inventories are disabled.

1. Outputs: Midday output, and hourly biogenic emissions from MEGAN
2. Midday and HEMCO outputs are used to determine isoprene to HCHO yield, after removing days with high smearing (Section 3.2.5)
3. HEMCO diagnostics are compared against top-down estimations of isoprene emissions (Section 3.3.1)

**Tropchem (altered MEGAN scaling factor)** Identical to standard tropchem except isoprene emissions are scaled to match multi-year monthly averaged top-down estimates.

1. Outputs: midday output, time series, daily averaged tracers
2. These are used to compare to campaign datasets after altering isoprene emissions (see Chapter 3)
3. They are also used to compare against satellite column HCHO as a sanity check on improving isoprene emissions

### 2.3.6.3 UCX vs tropchem

Here the model output is analysed before and after enabling the Universal tropospheric-stratospheric Chemistry eXtension (UCX). Both runs use  $2^\circ \times 2.5^\circ$  latitude by longitude; however, the UCX mechanism is run with 72 vertical levels from the surface to the top of the atmosphere (TOA  $\sim 0.1$  hPa), while the standard (tropchem) run uses 47 levels. The reduced vertical levels are created by lumping together sets of the higher resolved levels from around 70 hPa to the top of the atmosphere. For both runs the input parameters such as MEGAN emissions and GEOS-5 meteorological fields are identical.

Figure 2.12 shows an example of surface HCHO amounts, averaged over Jan and Feb, 2007, with and without the UCX mechanism enabled. Surface HCHO (first model level; up to  $\sim 100$  m) does not differ much between runs. The differences do not exceed

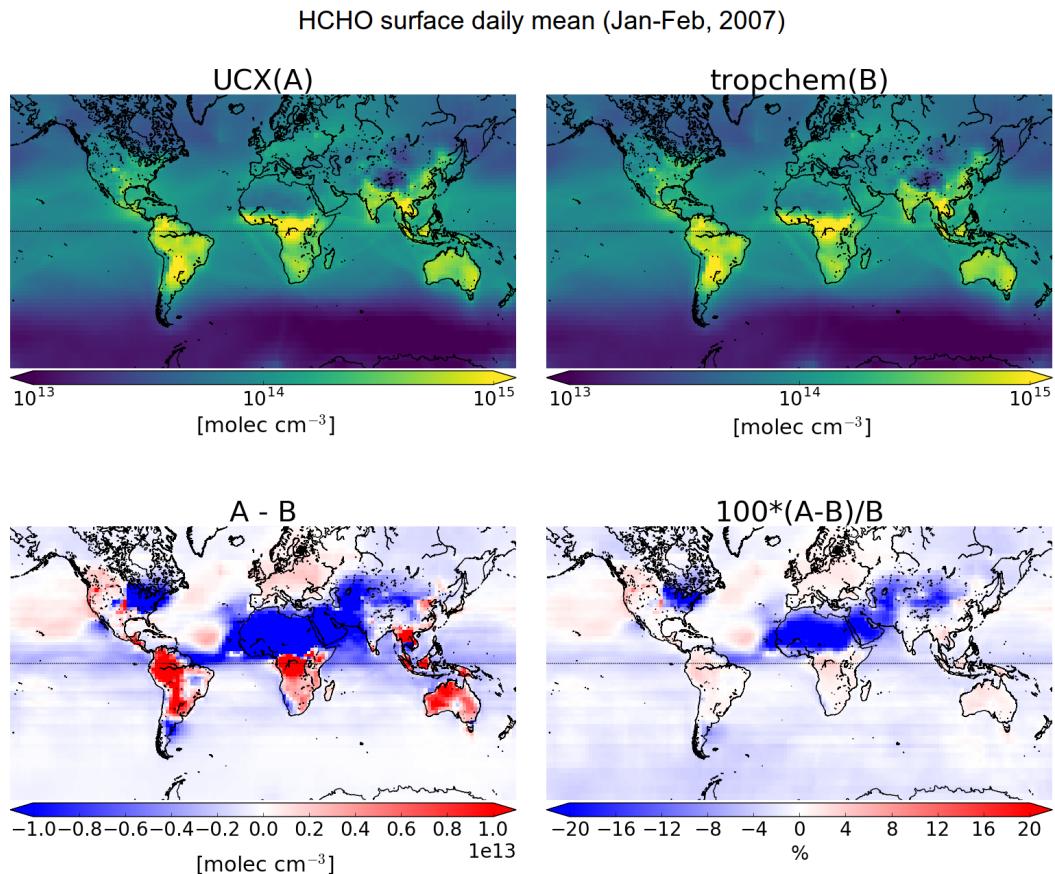


FIGURE 2.12: Surface HCHO simulated by GEOS-Chem with UCX (A: top left), and without UCX (B: top right), along with their absolute and relative differences (bottom left, right respectively). Amounts are the average of all times between 1 Jan and 28 Feb 2007.

3% over Australia, and absolute differences are minor (note the scale in A-B). The major notable difference occurs over northern Africa, where HCHO is around 20% lower in the UCX run (although absolute HCHO is relatively low in this region). Additionally a slight ( $< 8\%$ ) decrease in HCHO over the oceans can be seen. The comparison is repeated using total columns (instead of surface values) in Figure 2.13, showing that differences affecting HCHO between the model run are spread over the entire vertical column. Differences over Australia in total column HCHO are small, and these are the values used in this thesis.

Figure 2.14 shows the differences in surface isoprene concentrations over Australia, averaged over 1, Jan to 28, Feb, 2007. Here we start to see<sup>mh</sup> a higher relative difference in concentrations becomes apparent<sup>mh</sup>, although this is generally over the areas with lower absolute concentrations. Very little isoprene is seen away from the continents (4-5 orders of magnitude less), due to its short lifetime and lack of oceanic sources. Generally isoprene is 0-30% higher over mid to western Australia when the

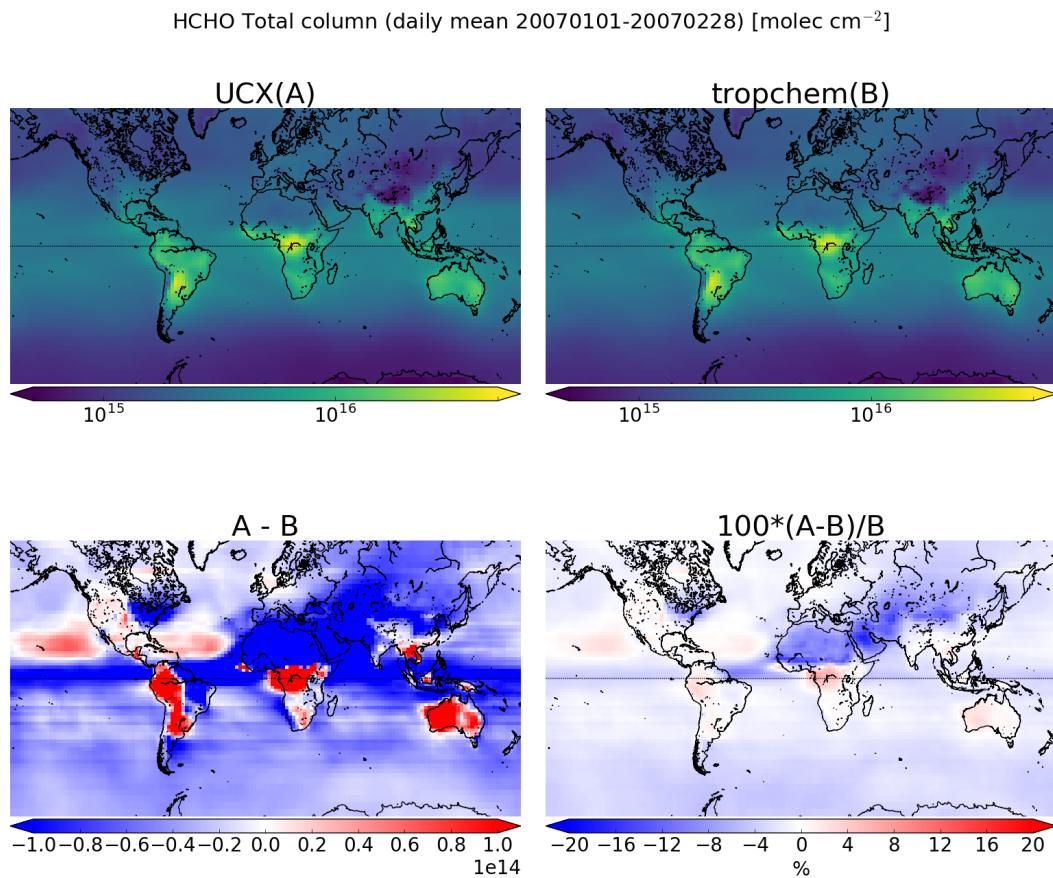


FIGURE 2.13: As Figure 2.12 but using total column amounts instead of surface concentrations.

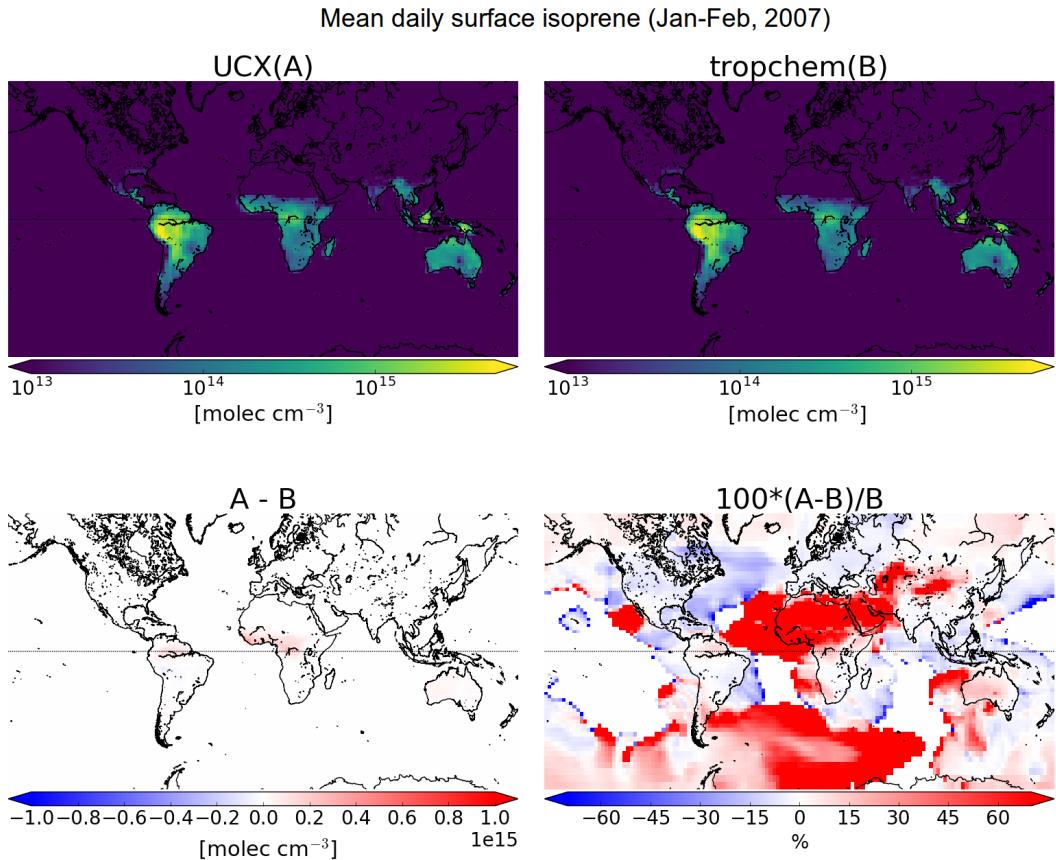


FIGURE 2.14: As Figure 2.12, except showing isoprene surface concentrations.

UCX mechanism is turned on; however, this increase is lower in the regions with high isoprene emissions (north-east to south-east coastline). This enhancement can be seen throughout the entire tropospheric column as shown by Figure 2.14. There is a greater effect in Africa and South America in the tropics, with high relative differences in many regions with low absolute amounts, but very small differences in locations where isoprene is strongly emitted.

The difference in isoprene between UCX and tropchem is likely caused by differences in the modelled radiation reaching the troposphere due to differences in simulated ozone in the stratosphere. Figure 2.16 shows the total column ozone between UCX and non-UCX run. This shows that UCX ozone is lower everywhere except for a thin band just north and south of the equator. With higher stratospheric ozone levels, less radiation would reach the troposphere in the tropchem run. This would slow photolysis limited reactions, such as the splitting of ozone that leads to OH production in the troposphere, in turn slowing the isoprene loss to OH reactions.

Overall the UCX mechanism and increased vertical resolution lead to slightly higher isoprene and HCHO, and slightly lower ozone over Australia. These differences are

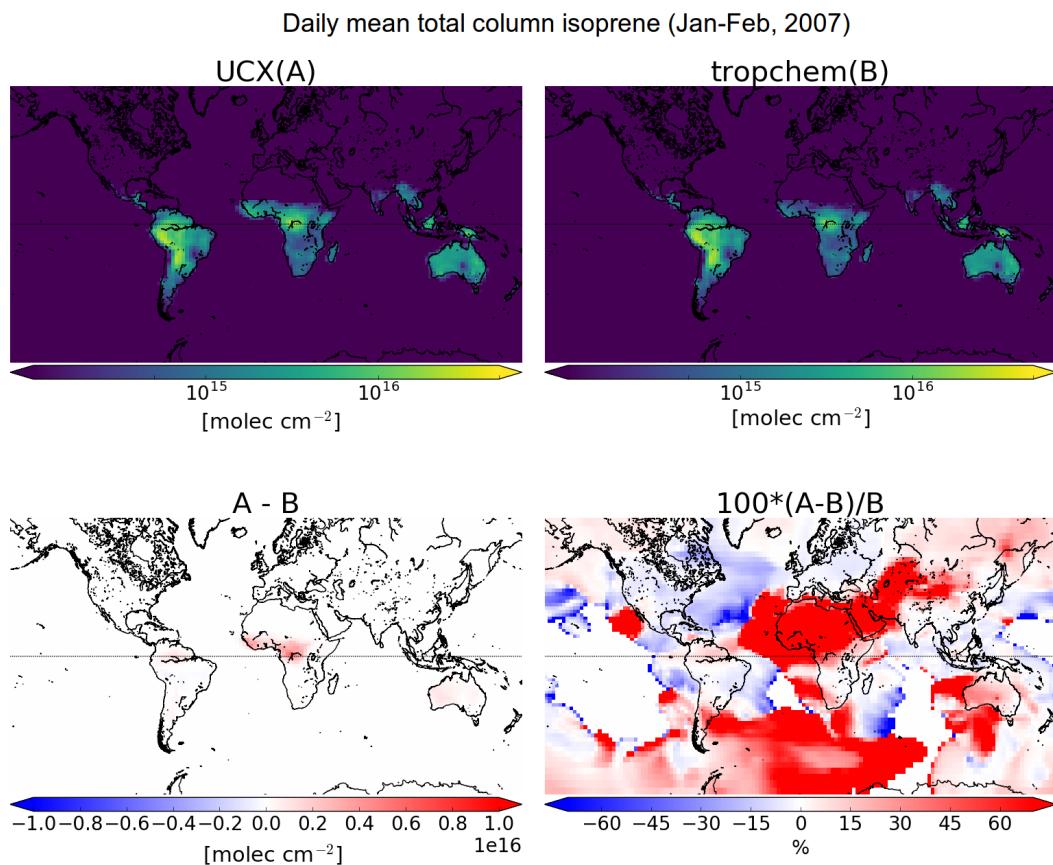


FIGURE 2.15: As Figure 2.14, except showing isoprene total column amounts.

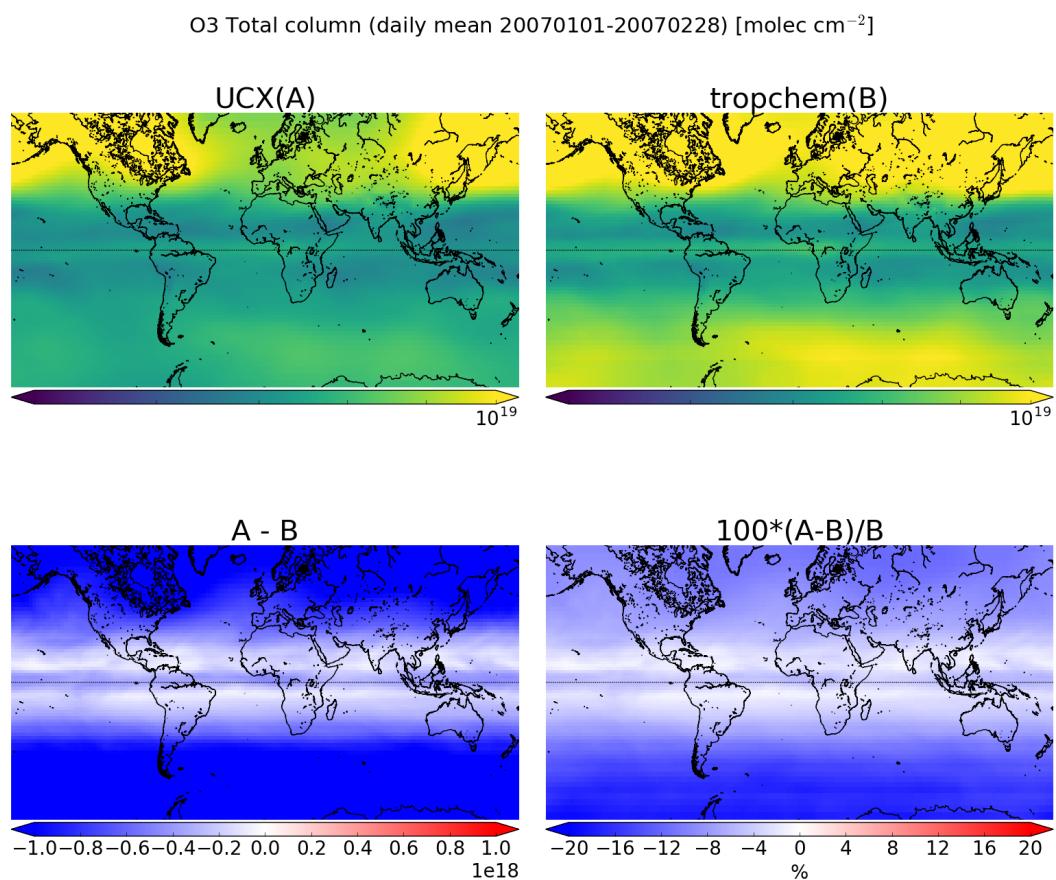


FIGURE 2.16: Total column ozone simulated by GEOS-Chem with UCX (A: top left), and without UCX (B: top right), along with their absolute and relative differences (bottom left, right respectively). Amounts are the average of all times between 1 Jan and 28 Feb 2007.

on the order of 0-10% and as shown later (see Section 3.4) are minimal compared to other uncertainties in both AMF calculation and emissions estimation. Running UCX requires roughly twice the computation (and real) hours, and outside of Chapter 4 the normal tropchem runs are used.

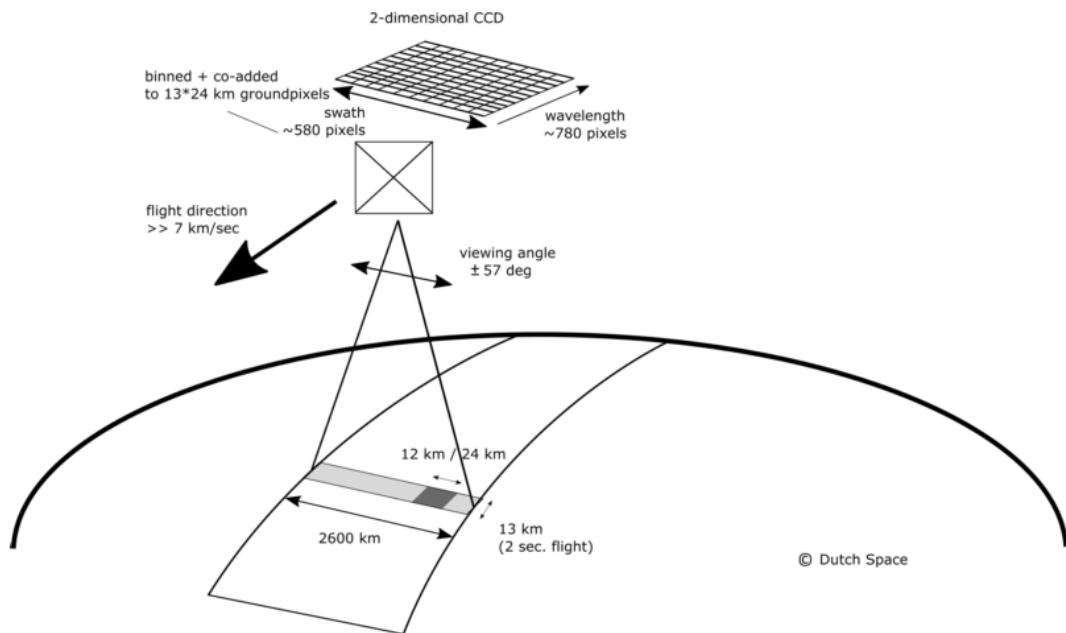
## 2.4 OMI satellite formaldehyde

One satellite product used extensively in this thesis is OMHCHO from NASA's Earth Observing System's Aura satellite. Aura orbits the earth in a polar sun-synchronous pattern, circling the earth on a (geometric) plane coincident with the sun and the poles. Aura houses the Ozone Monitoring Instrument (OMI), a near-UV/Visible Charged Coupled Device (CCD) spectrometer. The OMI instrument onboard Aura has been active since July 2005. It records spectra from 264-504 nm using an array of 60 detectors with moderate resolution (0.4-0.6 nm). This band of wavelengths allows measurements of trace gases (among other quantities) and the formaldehyde product is detailed here.

From here onward the word pixel is used to describe one data point retrieved by OMI. Each pixel includes a latitude and longitude within OMI's data product. Figure 2.17 shows the details of OMI's detector array and measurement resolutions. OMI measurements occur from right to left on a band covering a viewing angle of  $115^\circ$ , resulting in swaths of around 2600 km, with pixel sizes from  $13 \times 24 \text{ km}^2$  at nadir to  $26 \times 135 \text{ km}^2$  at the swath edges (Gonzalez Abad et al. 2015). The swaths cover the Earth daily, both on the light and dark side of the planet, but only daytime measurements provide useful near-UV/Visible information. While satellite measurements can only be used during daytime hours, HCHO lifetimes are sufficiently short that any nighttime chemistry will not affect midday observations (Wolfe et al. 2016).

The latest OMHCHO algorithm uses a shape factor determined from GEOS-Chem (V9) using 47 vertical levels at monthly temporal resolution and  $2^\circ \times 2.5^\circ$  latitude by longitude horizontal resolution (Gonzalez Abad et al. 2015). The GEOS-Chem model has been substantially updated since then. In this thesis a more recent version V10.01 is used to recalculate the vertical column HCHO (details are shown in Section 2.6).

OMI uses a Differential Optical Absorption Spectroscopy (DOAS) based technique to retrieve HCHO along the path of light that reaches the satellite instrument. The first step is to determine how much HCHO is in the path of light between the sun and detector, which is done by applying a forward radiative transfer model (RTM) (see Section 2.4.2) in order to determine the radiative properties of a trace gas at various altitudes. The forward RTM used for satellite data products also involve functions representing extinction from Mie and Rayleigh scattering, and the effect of these on spectra. Rayleigh and Mie scattering (two kinds of particle effects on radiation passing through a medium) must be accounted for in the UV-VIS region of the spectrum. These RTM are also required to account for (often estimated) atmospheric parameters such as albedo. The next step is to transform the calculated amounts along the non-vertical light path into vertical column amounts. This is done by applying an AMF. In the absence of atmospheric scattering a simple geometric AMF (a scaling factor using the projection of the slant column onto a vertical axis) can be defined as a function of the



Channel	Wavelength range	Spectral resolution	Spectral sampling	Ground pixel size
UV1	264–311 nm	0.63 nm = 1.9 px	0.33 nm px <sup>-1</sup>	13 × 48 km
UV2	307–383 nm	0.42 nm = 3.0 px	0.14 nm px <sup>-1</sup>	13 × 24 km
VIS	349–504 nm	0.63 nm = 3.0 px	0.21 nm px <sup>-1</sup>	13 × 24 km

FIGURE 2.17: Here is an impression of the OMI scanning swathes as the housing satellite sweeps around the Earth. The charge-coupled device (CCD) has two dimensions: wavelengths and cross-tracks. The table shows the optical properties for OMI's three channels. Reproduced from Schenkeveld et al. (2017).

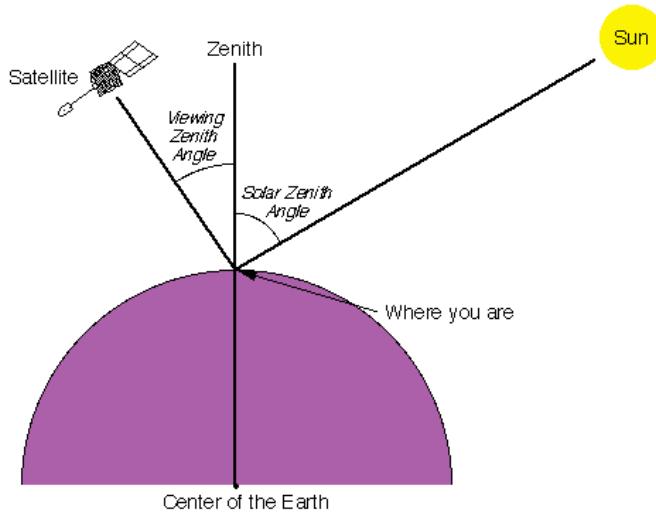


FIGURE 2.18: Solar and viewing zenith angles, reproduced from <http://sacs.aeronomie.be/info/sza.php>.

solar zenith angle. The solar zenith angle ( $\theta_s$ ) and the satellite viewing angle ( $\theta_v$ ) are shown in image 2.18.

In this thesis I use<sup>mh</sup> the NASA OMHCHOv003 data product (Gonzalez Abad et al. 2015) is used<sup>mh</sup>, with HCHO determined using the spectral window 328.5 nm–356.5 nm. The algorithm used is based on direct fitting of radiances, and accounts for competing absorbers, under-sampling, and Ring effects (these affect measured absorption lines). Slant columns ( $\Omega_S$ ) are determined from the spectra differential with respect to radiance reference column over the remote Pacific. The full method details for slant column retrieval by OMI are outlined in the technical document ([https://docserver.gesdisc.eosdis.nasa.gov/repository/Mission/OMI/3.3\\_ScienceDataProductDocumentation/3.3.4\\_ProductGenerationAlgorithm/ATBD-OMI-04.pdf](https://docserver.gesdisc.eosdis.nasa.gov/repository/Mission/OMI/3.3_ScienceDataProductDocumentation/3.3.4_ProductGenerationAlgorithm/ATBD-OMI-04.pdf)). Slant columns range from  $\sim 4 \times 10^{15}$  to  $\sim 6 \times 10^{16}$  molec cm<sup>-2</sup>, with uncertainties from 30% (larger columns) to over 100% (smaller columns) (Gonzalez Abad et al. 2015).

### 2.4.1 Pixel filtering

This thesis uses the level two OMHCHO product swath output from the NASA earth data web portal. OMHCHO level two data includes 14-15 daily swaths of measurements. Each swath contains roughly  $9 \times 10^4$  pixels, each of which includes latitude, longitude, vertical column HCHO, along with all the ancillary data required to calculate the vertical column and several data quality metrics. The OMHCHO dataset has a quality flag that can be used to remove unlikely or poor satellite measurements. The states represented by this quality flag are shown in Table 2.3 that is reproduced here from Kurosu and Chance (2014). First all *good* pixels (those with QA flag equal to 0) are read into a long list (roughly 1 million per day). Filtering bad or missing measurement pixels is performed prior to any other filtering. This includes the data points affected by the row anomaly (see [https://www.cfa.harvard.edu/atmosphere/Instruments/OMI/PGEReleases/READMEs/OMHCHO\\_README\\_v3.0.pdf](https://www.cfa.harvard.edu/atmosphere/Instruments/OMI/PGEReleases/READMEs/OMHCHO_README_v3.0.pdf)). This anomaly affects radiance

TABLE 2.3: OMI quality flag values from Kurosu and Chance (2014)

<b>Value</b>	<b>Classification</b>	<b>Rational</b>
0	Good	Column value present and passes all quality checks; data may be used with confidence.
1	Suspect	Caution advised because one or more of the following conditions are present: <ul style="list-style-type: none"> <li>• Fit convergence flag is <math>&lt; 300</math> but <math>&gt; 0</math>: Convergence at noise level</li> <li>• Column <math>+2\sigma</math> uncertainty <math>&lt; 0 &lt;</math> Column <math>+3\sigma</math> uncertainty</li> <li>• Absolute column value <math>&gt;</math> Maximum column amount (<math>1e19</math> molec <math>cm^{-2}</math>)</li> </ul>
2	Bad	Avoid using as one of the following conditions are present: <ul style="list-style-type: none"> <li>• Fit convergence flag is <math>&lt; 0</math> : No convergence, abnormal termination</li> <li>• Column <math>+3\sigma</math> uncertainty <math>&lt; 0</math></li> </ul>
$< 0$	Missing	No column values have been computed; entries are missing

data at particular viewing angles, corresponding to a row on the CCD detectors, and is dynamic over time (Huang et al. 2018). The slant columns affected are flagged and removed before any further processing.

Approximately every 90 minutes, the Aura satellite sweeps over the sun lit side of the planet and makes around 90000 measurements, of which 50000 - 80000 pixels are classified as good. Each pixel contains several important pieces of data that are needed for recalculation of the HCHO vertical column: the total column of HCHO ( $\Omega$ ; molec  $cm^{-2}$ ), cloud fraction, associated shape factor, AMF, geometric AMF, scattering weights and their vertical altitudes (hPa), viewing zenith angle, solar zenith angle, latitude, longitude, OMI sensor track, main data quality flag, cross track flag, and total column uncertainty. All of these data are needed in order to reconstruct the total vertical column using a new a priori estimate of the vertical profile in lieu of that provided by NASA. This reconstruction is required so that perceived bias between satellite measurements and model outputs is not due to a priori information, but rather what the satellite is producing.

After being filtered for data quality, pixels are filtered by solar zenith angle (SZA) and latitude, similarly to other works (e.g., Marais et al. 2012; Barkley et al. 2013; Zhu et al. 2014b; Bauwens et al. 2016; Zhu et al. 2016). Satellite measurements poleward of  $60^\circ$  north or south are removed as well as measurements with SZA greater than  $60^\circ$ . Measurements with high SZA are unlikely to have much information near the surface, and often have high uncertainty in total column retrievals (e.g., Stone et al. 2015). Pixels with cloud fraction greater than 40% are removed after determining the

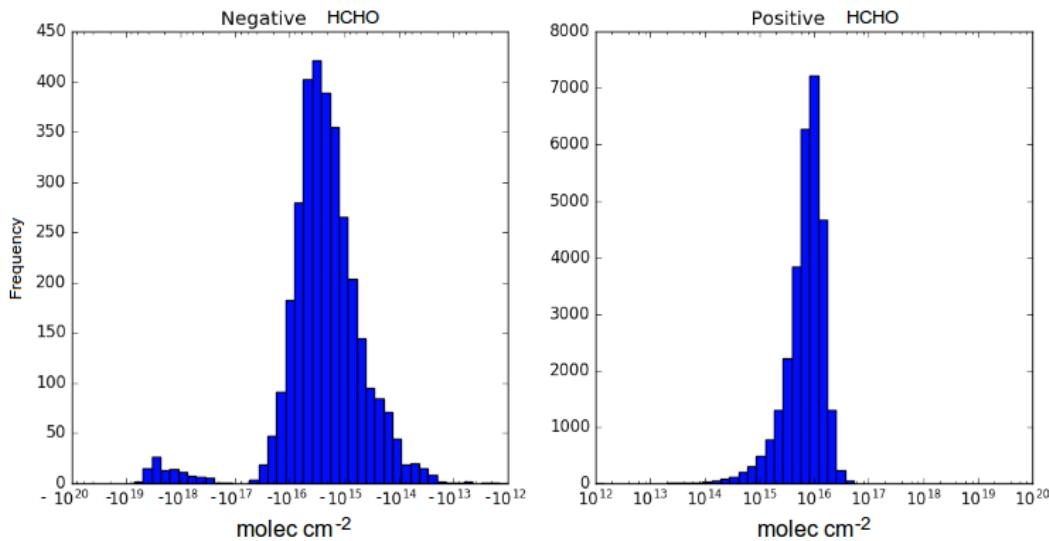


FIGURE 2.19: HCHO Column amount histograms for a subset of OMI swaths over Australia on the 18th of March 2013. Negative entries are shown in the left panel, positive in the right. Note the different scale between negative and positive panels.

reference sector correction (Section 2.6.5), as in Gonzalez Abad et al. (2015) and De Smedt et al. (2015). This removes around 30% of the pixels that remain after filtering out the bad or missing data.

Due to noise and the differential measurement technique, small negative columns are present in the satellite product that are not removed so as not to introduce a bias. However, some very large negative values are removed, as these represent an unflagged error in the retrieval process. Unlikely high positive measurements are also removed, leaving only measurements within the range  $-0.5 \times 10^{16}$  to  $1 \times 10^{17}$  molec  $\text{cm}^{-2}$ , as is performed by Zhu et al. (2016). This filter is required due to currently unexplained large negative values which occur in the OMI HCHO product increasingly over time. Figure 2.19 shows how unfiltered HCHO columns are affected by a small set of highly negative values that heavily affect the mean column amount over any region. The highly negative values can be seen around  $\Omega = -10^{19}$  molecules  $\text{cm}^{-2}$ .

#### 2.4.2 Air mass factor (AMF)

To convert the trace gas profile from a reflected solar radiance column (slanted along the light path) into a purely vertical column requires calculations of an air mass factor (AMF). In satellite data, the AMF is typically a scalar value for each horizontal grid point that gives the ratio of the total vertical column density to the total slant column density. An AMF characterises measurement sensitivity to a trace gas at various altitudes (Palmer et al. 2001). This value is unique for each trace gas under consideration (Palmer et al. 2001; Millet et al. 2006). Lorente et al. (2017) show that AMF calculations can be the largest source of uncertainty in satellite measurements. Calculation of the

AMF is important as it is multiplied against the estimated slant columns in order to give vertical column amounts.

DOAS column retrievals are an integration of a trace gas over the instrument's viewing path. In order to convert this total to a vertically distributed column a few assumptions and estimates are required. The initial (*a priori*) vertical profile of a trace gas is assumed or estimated via a CTM, while its scattering and radiative properties are calculated at prescribed altitudes using an RTM. These properties are combined to determine the AMF. Different models are used for different satellite AMFs. For example GOME-2 products use the LInearized Discrete Ordinate Radiative Transfer (LIDORT) RTM with IMAGESv2 CTM, and OMI products use LIDORT combined with the GEOS-Chem CTM (Chance 2002; Gonzalez Abad et al. 2015). Lamsal et al. (2014) recommends that when comparing satellite data to models, the AMF should first be recalculated using the model as an *a priori*. This is in order to remove any *a priori* bias between model and satellite columns. For this reason the OMHCHO AMF are recalculated in this thesis using GEOS-Chem to create the *a priori* profiles and LIDORT to calculate scattering weights.

LIDORT is a model of LInearized Discrete Ordinate Radiative Transfer, used to determine backscatter intensities and weighting functions at arbitrary elevation angles (Spurr, Kurosu, and Chance 2001). The model solves radiative transfer equations and can be used to determine various atmospheric column measurement attributes such as optical depth, ring effects, and scattering. These radiative properties (or at least estimates thereof) are required when measuring trace gases in the atmosphere through a long path such as seen by satellites (e.g., Palmer et al. 2001; Martin et al. 2002; De Smedt et al. 2015; Gonzalez Abad et al. 2015).

Related to the AMF is the averaging kernel (AK), which is used to handle instrument measurements that are sensitive to gas concentrations at different altitudes through the atmosphere. DOAS methods can be heavily influenced by the initial estimates of a trace gas profile (*the a priori*) that is often produced by modelling, so when comparing models of these trace gases to satellite measurements extra care needs to be taken to avoid introducing bias from differing *a priori* assumptions. One way to remove these *a priori* influences is through the satellite AK (or AMF), which takes into account the vertical profile of the modelled trace gas and instrument sensitivity to the trace gas (Eskes and Boersma 2003; Palmer et al. 2001). This process is called deconvolution ( $\Omega = AK \times VC_{satellite} + (I - AK) \times VC_{apriori}$ ) of the AK of the satellite instrument. The AK represents sensitivities to each species at multiple altitudes through the atmosphere and in the case of OMI, can be approximated from the scattering weights ( $\omega(z)$ ) function as follows:

$$AK(z) = \frac{\omega(z)}{AMF} \quad (2.7)$$

(Gonzalez Abad et al. 2015).

### 2.4.3 Uncertainty

The main sources of error in satellite retrievals of HCHO are instrument detection sensitivities and calculation of the AMF (the factor calculated to convert slanted light path concentrations into a vertical profile) (Millet et al. 2006). Calculations of the AMF performed by different groups tend to agree fairly well, as long as the *a priori* and

ancillary data are similar. Large differences can occur depending on the a priori vertical profile, trace gas concentrations, and cloud properties (Lorente et al. 2017). Choice of radiative transfer model and interpolation operations have a relatively small affect compared to the assumed state of the atmosphere, with high structural uncertainty introduced at this stage of AMF calculation (Lorente et al. 2017).

Uncertainty in the OMI satellite instrument is calculated by the Smithsonian Astrophysical Observatory (SAO) group using the uncertainty in backscattered radiation retrievals (Gonzalez Abad et al. 2015; Abad et al. 2016). Uncertainty in a single pixel for OMHCCHO is roughly the same magnitude as HCHO background levels. Each pixel has  $\sim 2 \times 10^{14}$  molec cm $^{-2}$  uncertainty, which is five times higher than GOME. However, there are  $\sim 100 - 200$  times as many measurements allowing a greater reduction of uncertainty with averaging. This is due to the smaller footprint and better temporal resolution of OMI (Chance 2002; Millet et al. 2008). The finer nadir resolution of OMI (13 by 24 km $^2$ ) compared to other satellites also reduces cloud influence (Millet et al. 2006; Millet et al. 2008). The top row in Figure 2.20 shows OMI HCHO columns binned to at  $0.25^\circ \times 0.3125^\circ$  longitude by latitude averaged over one day and one month (with and without filtering). Row two shows uncertainty of the satellite data after averaging. It is clear that one day of satellite data is too uncertain when binned at  $0.25^\circ \times 0.3125^\circ$  horizontal resolution; however, after a month (with or without filtering) the uncertainties become manageable. If it is assumed that we assume<sup>mh</sup> the uncertainty is random error, and not bias introduced through calculation techniques, then this uncertainty can be reduced we are able to reduce the uncertainty<sup>mh</sup> through averaging. Uncertainty in satellite recalculations related to estimating isoprene emissions is analysed in Section 3.4.3.

Recently Schenkeveld et al. (2017) analysed the instrument performance over the life time (approximately 2004 to 2015) of OMI and found irradiance degradation of 3-8%, changed radiances of 1-2%, and a stable wavelength calibration within 0.005-0.020 nm. These changes are measured excluding the row anomaly effect, which has been relatively stable since 2011, although it is still growing and remains the most serious concern. Their analysis concludes that the satellite data is still of high quality and will deliver useful information for 5-10 more years, with radiance only changing by 1-2% outside of row anomaly impacted areas. An analysis of the row anomaly by Huang et al. (2018) states that measurements remain suitable for scientific use, with recommendation for further evaluation. The row anomaly began in June 2007, with some cross-track rows seemingly blocked. The most likely cause is some instrument insulation partially obscuring the radiance port (Schenkeveld et al. 2017). In this thesis pixels potentially affected by the row anomaly are removed prior to calculation or analysis.

In satellite HCHO products, concentrations over the remote Pacific ocean are sometimes used to analyse faulty instrument readings. This is due to the expected invariance of HCHO over this region. For instance GOME (an instrument that measures trace gases onboard the ERS-2) corrects for an instrument artefact using modelled HCHO over the remote Pacific (Shim et al. 2005). OMI HCHO products use a similar technique to account for sensor plate drift and changing bromine sensitivity (Gonzalez Abad et al. 2015). In this thesis a background uncertainty estimation is performed based on differences between satellite measurements and monthly averaged modelled

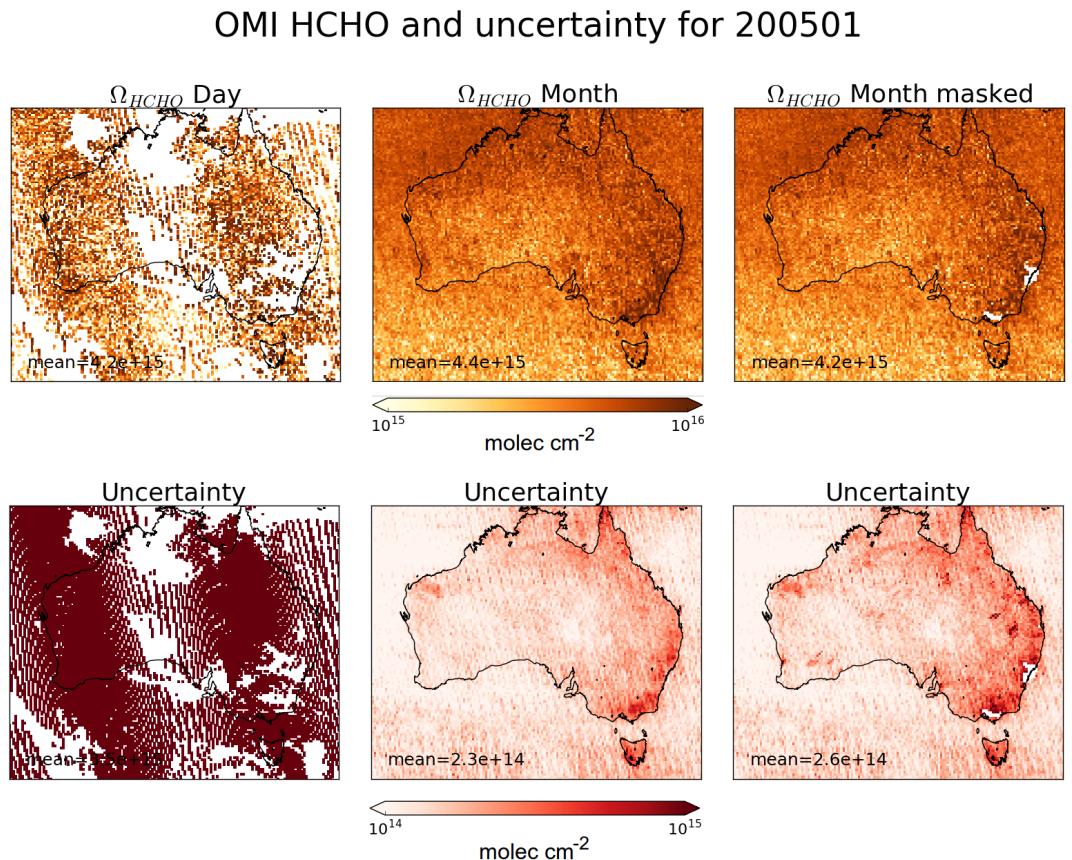


FIGURE 2.20: Top row shows  $0.25^\circ \times 0.3125^\circ$  binned OMI HCHO columns with one day, one month, and one month with non-biogenic masking applied from left to right respectively. Bottom row shows the uncertainty for each grid square after averaging.

HCHO over the remote Pacific ocean (see Section 3.4.3).

The OMI dataset may suffer from bias; however, in order to determine any bias an independent dataset is required, and this does not exist over Australia. In this thesis, HCHO columns (pixels) with cloud fractions over 40% are filtered as done in Palmer et al. (2001), which may introduce a clear-sky bias to any monthly averages. This is due to HCHO being lower on unrecorded cloudy days. This bias has been measured as a 13% positive monthly mean bias (Palmer et al. 2001; Surl, Palmer, and Abad 2018). For many places the tropospheric column HCHO measured by satellite is biased low. Zhu et al. (2016) examine OMHCHO and show a bias of 37% over south-east USA when compared against a campaign of aircraft observations (SEAC<sup>4</sup>RS). De Smedt et al. (2015) also found OMI and GOME2 observations were 20 - 40% lower than ground based vertical profiles, and Barkley et al. (2013) determine OMI to be 37% low compared with aircraft measurements over Guyana. Millet et al. (2006) examine OMI HCHO columns over North America and determine overall uncertainty to be 40%, with most of this coming from cloud interference. Millet et al. (2008) shows that there also exists some latitude based bias, as well as a systematic offset between the OMI and GOME instruments. This does not appear to be due to the different overpass times of the two instruments.

AMF calculation often dominates the total uncertainty in satellite retrievals, especially in polluted regions (Lorente et al. 2017). In scenarios where HCHO is enhanced in the lower troposphere, AMF calculation is the largest uncertainty in satellite measurements. In polluted environments the structural uncertainty is estimated at 42 %, or 31 % over unpolluted environments (Lorente et al. 2017). Another impact often not included in uncertainty calculations is the structural uncertainty of retrieval methods. The structural uncertainty of AMF calculation approaches used by different retrieval groups comes from how the AMF is calculated, rather than uncertainty in the calculation components. The importance of a priori and ancillary data (such as surface albedo and cloud top height) sharply affects the structural uncertainty (Lorente et al. 2017). In this thesis the AMF uncertainty is assumed to be 30%, as concentrations of HCHO and pollution are relatively low over Australia.

## 2.5 Calculating new AMF

One of the uses of GEOS-Chem in this thesis is to update the a priori HCHO column used in satellite measurements. The a priori is used when deriving the air mass factor (*AMF*), which is needed to transform satellite slant columns (*SC*) into vertical columns ( $\Omega$  in molecules  $\text{cm}^{-2}$ ): The AMF is the ratio of *SC* to the vertical column

$$\text{AMF} = \frac{\text{SC}}{\Omega} = \frac{\tau_s}{\tau_v} \quad (2.8)$$

with  $\tau$  being the optical depth or thickness of the absorber through the slant (*s*) or vertical (*v*) path of light.

The OMI instrument records spectra of light that enter the viewing lens on the Aura satellite. The spectra provide backscattered intensity ( $I_B$ ) at various wavelengths (see Section 2.4), with the light source ( $I_{B_0}$ ) being the sun. Using the log of Beer's law

(Equation 1.8) gives we get<sup>mh</sup>

$$\tau_s = \ln I_{B_0} - \ln I_B$$

which can be substituted into Equation 2.8 to give an expression for the AMF that includes scattering:

$$AMF = \frac{\ln I_{B_0} - \ln I_B}{\tau_v} \quad (2.9)$$

We use<sup>mh</sup>  $\nabla I = I_B - I_{B_0}$  is used<sup>mh</sup> to represent the change in intensity due to the absorber. For optically thin absorption,  $\nabla I / I_B \ll 1$ , which allows one to, and we can<sup>mh</sup> use:

$$AMF = \frac{\ln \left(1 - \frac{\nabla I}{I_B}\right)}{\tau_v} \approx \frac{-\frac{\nabla I}{I_B}}{\tau_v} \quad (2.10)$$

This is due to the logarithmic property  $\ln(1 - x) \approx -x$  for  $x \ll 1$ .  $\nabla I$  can also be expressed as the integral of the absorption slices over optical depth increments:

$$\nabla I = \int_0^{\tau_v} \frac{\partial I_B}{\partial \tau} d\tau \frac{\nabla I}{I_B} = \int_0^{\tau_v} \frac{\partial \ln I_B}{\partial \tau} d\tau$$

which can be placed into Equation 2.10 leading to

$$AMF \approx \frac{-1}{\tau_v} \int_0^{\tau_v} \frac{\partial \ln I_B}{\partial \tau} d\tau$$

Conversion of  $d\tau$  to the satellite path is performed We can then convert  $d\tau$  to our path<sup>mh</sup> using Equation 1.10 leading to

$$AMF = \frac{-1}{\tau_v} \int_0^{\infty} \frac{\partial \ln I_B}{\partial \tau} \alpha(z) \eta(z) dz \quad (2.11)$$

where  $\alpha(z)$  and  $\eta(z)$  represent absorption cross section in  $\text{m}^2 \text{ molecule}^{-1}$ , and number density in molecules  $\text{m}^{-3}$  respectively. This uses the attenuation cross section relationship to optical depth (see Section 1.4.2.2).

To represent an average cross section weighted by the absorbing species' vertical distribution, the effective cross section ( $\alpha_e$ ) is used. This is to account for temperature and pressure dependence of  $\alpha(z)$ , and is defined as:

$$\begin{aligned} \alpha_e &= \frac{1}{\Omega_v} \int_0^{\infty} \alpha(z) \eta(z) dz \\ &= \frac{\tau_v}{\Omega_v} \end{aligned}$$

Then replacing the  $\tau_v$  in Equation 2.11 provides we obtain<sup>mh</sup>.

$$AMF = - \int_0^{\infty} \frac{\partial \ln I_B}{\partial \tau} \frac{\alpha(z)}{\alpha_e} \frac{\eta(z)}{\Omega_v} dz \quad (2.12)$$

Often the integrand of this AMF formula (Equation 2.12) is broken apart into two defining terms: the scattering weights  $\omega(z)$  and the shape factor  $S(z)$ , described here:

- $\omega$  The scattering weights describing sensitivity of the backscattered spectrum to the abundance of an absorber at altitude z:

$$\omega(z) = -\frac{1}{AMF_G} \frac{\alpha(z)}{\alpha_e} \frac{\partial \ln I_B}{\partial \tau} \quad (2.13)$$

It is worth noting that in the OMI satellite product, the provided  $\omega(z)$  term does not include the  $\frac{1}{AMF_G}$  term and the calculations that follow therefore do not include this term when utilising the provided  $\omega$ . This is not noted in any of the papers that recalculate the AMF from the OMI product, due to them recalculating the  $\omega$  term themselves with a radiative transfer model such as LIDORT.

- S the shape factor describes the profile of an absorber ( $\eta(z)$ ) normalised by its total vertical column amount ( $\Omega_v$ ):

$$S(z) = \frac{\eta(z)}{\Omega_v} \quad (2.14)$$

Plugging Equations 2.13 and 2.14 into Equation 2.12 gives ~~us~~<sup>mh</sup>:

$$AMF = AMF_G \int_0^\infty \omega(z) S(z) dz \quad (2.15)$$

Since ~~this thesis uses we are using~~<sup>mh</sup> the  $\omega$  provided by OMI, the  $AMF_G$  term is removed from this calculation as it is not part Equation 2.13 leading to

$$AMF = \int_0^\infty \omega(z) S(z) dz \quad (2.16)$$

Additionally the AMF can be determined using the sigma ( $\sigma$ ) coordinate system, which denotes altitude as going from 1 (ground level) to 0 (TOA) (see Section 2.2). A conversion to the  $\sigma$  vertical coordinate is performed using  $p = \sigma(p_S - p_T) + p_T$ , where  $p_T$  is pressure at the top of the atmosphere and  $p_S$  is surface pressure.  $S_\sigma$  is a dimensionless normalised shape factor on the  $\sigma$  coordinate system. In the sigma coordinate system ~~we calculated~~<sup>mh</sup> the shape factor as defined in Palmer et al. (2001) ~~is calculated as follows~~<sup>mh</sup>:

$$S_\sigma(\sigma) = \frac{\Omega_a}{\Omega_v} C_{HCHO}(\sigma) \quad (2.17)$$

where  $\Omega_a$  is the vertical column of air from the surface to the top of the atmosphere and  $C_{HCHO}(\sigma)$  is the mixing ratio of HCHO. Then one ~~We~~<sup>mh</sup> can integrate over the sigma coordinates using the hydrostatic relation  $p = -\rho_a g z$ , with  $\rho_a$  being air density, and  $g$  the gravitational acceleration:

$$\begin{aligned} \rho_a g z &= \sigma(p_S - p_T) + p_T \\ d\sigma &= -\frac{\rho_a g}{p_S - p_T} dz \end{aligned}$$

Substitution into 2.16 gives AMF using the sigma coordinates:

$$AMF = \int_0^1 w(\sigma) S_\sigma(\sigma) d\sigma \quad (2.18)$$

## 2.6 Recalculation of OMI HCHO

OMI HCHO vertical columns are calculated using modelled a priori HCHO profiles (see Section 2.4). When comparing satellite measurements against models it is important to recognise the impact of this a priori on the total column values. This is complicated by how OMI is differently sensitive to HCHO (and other trace gases) vertically throughout the atmosphere. When comparing OMI vertical columns ( $\Omega_{OMI}$ ) to GEOS-Chem ( $\Omega_{GC}$ ), the satellite AMF needs to be recalculated using GEOS-Chem modelled vertical gas profiles as the a priori. Without performing this step a bias between modelled and measured total column values may be due to the a priori rather than chemistry or measurements (Palmer et al. 2001; Lamsal et al. 2014).

To recalculate vertical columns two components are calculated: an AMF and a reference sector correction. Here, two new AMFs are calculated, both using GEOS-Chem HCHO profiles as the new a priori. The first ( $AMF_{GC}$ ) recalculates the satellite shape factor, but uses the original satellite scattering weights. The second ( $AMF_{PP}$ ) recalculates both the satellite shape factor and the scattering weights.  $AMF_{PP}$  is created using code initially written by Professor P. Palmer.

In addition to a new AMF, a reference sector correction is determined using the method described in Abad et al. (2016). This correction has also been applied to the original vertical column in the OMHCHO product, so the column is available with and without this correction. The reference sector correction is unique for each of the 60 measurement tracks used by OMI. The correction is calculated for each of the two new vertical columns, and applied to each pixel to create the corrected vertical columns. The end product is three sets of corrected vertical columns: the original, one based on new shape factors, and one with new shape factors and scattering weights.

### 2.6.1 Outline

An outline in computational order of what takes place when recalculating  $\Omega$  from OMI follows:

1. Section 2.6.2 describes how GEOS-Chem satellite overpass output is used to create new shape factors ( $S_z$  and  $S_\sigma$ ).
  - (a) Vertical pressure edges and geometric midpoints are determined, along with altitudes ( $z$ ), and box heights  $H$ .
  - (b) Number density and mixing ratio of HCHO ( $n_{HCHO}$ ,  $C_{HCHO}$  respectively) are taken or created from model outputs HCHO (ppb), air density (molec  $\text{cm}^{-3}$ ), and box heights (m).
  - (c) Total column HCHO from GEOS-Chem ( $\Omega_{GC}$ ) is calculated

$$\Omega_{GC} = \Sigma_z (n_{HCHO}(z) \times H(z))$$

along with total column air ( $\Omega_A$ , calculated similarly).

- (d) The shape factor  $S_z(z)$  is calculated for each altitude

$$S_z(z) = n_{HCHO}(z) / \Omega_{HCHO}$$

- (e) Sigma coordinates are created from pressures (see Section 2.2, Equation 2.1).
- (f)  $S_\sigma(z)$  is calculated on each altitude:

$$S_\sigma(z) = C_{HCHO}(z) \times \frac{\Omega_A}{\Omega_{HCHO}}$$

2. For each pixel, new AMF ( $AMF_{GC}$  and  $AMF_{PP}$ ) is created using the GEOS-Chem shape factors and scattering weights from the satellite product or from the Palmer code in Section 2.6.3:

- (a) Satellite pixels (SC, scattering weights ( $\omega(z)$ ), pressure levels, latitude and longitude) are read from the OMHCHO dataset.
- (b) scattering weights ( $\omega$ ) are interpolated onto the same vertical dimensions ( $z$  and  $\sigma$ ) as the shape factors.
- (c) Integration (approximated using rectangular method) is performed along the vertical dimension to calculate the new AMF on both coordinate systems:

$$AMF_z = \Sigma_z (\omega(z) \times S_z(z) \times H(z)) \quad (2.19)$$

$$AMF_\sigma = \Sigma_\sigma (\omega(\sigma) \times S_\sigma(\sigma) \times d\sigma) \quad (2.20)$$

These two AMFs represent the same thing using different coordinates, and either one can be used as the  $AMF_{GC}$ .

- (d) Code from Dr. Surl and Professor Palmer is used to recalculate both the shape factor and scattering weights to produce  $AMF_{PP}$ .
3. Section 2.6.4 describes how the original AMF,  $AMF_{PP}$ , and  $AMF_{GC}$  are used to determine the new vertical columns:  $\Omega = SC / AMF$ .
  4. A reference sector correction (RSC) is defined each day using these AMFs along with modelled HCHO over the remote Pacific in Section 2.6.5:
    - (a) GEOS-Chem midday output ( $\Omega_{GC}$  from  $140^\circ W$  to  $160^\circ W$ ) is averaged monthly and longitudinally to provide modelled vertical columns over the reference sector.
    - (b) These vertical columns ( $\Omega_{GC}$ ) are turned into modelled slant columns by multiplication with the AMF over the reference sector:

$$SC_{GC} = \Omega_{GC} \times AMF$$

- (c) For each satellite pixel in the reference sector, a correction ( $corr$ ) is calculated as the measured  $SC$  minus the modelled slant column at the nearest

latitude:

$$\text{corr}[lat, track] = SC[lat, track] - \Omega_{GC}[lat] \times AMF$$

- (d) These corrections are binned by satellite detector (track: 1-60), and latitude ( $0.36^\circ$ ; 500 latitudes from  $90^\circ\text{S}$  to  $90^\circ\text{N}$ ).
  - (e) The median entry of each bin is determined and this forms the  $RSC[lat, track]$  (e.g., Figure 2.24).
5. Corrected vertical columns  $VCC$  are determined using  $VCC = (SC - RSC[lat, track]) / AMF$  for each measured  $SC$  and using each  $AMF$ , with the  $RSC$  linearly interpolated to the latitude of the satellite pixel (see Section 2.6.6).
  6. The  $VCC$  (along with most of the pixel and GEOS-Chem data) are binned onto a  $0.25^\circ \times 0.3125^\circ$  grid along with other information (see Section 2.6.7).

Each of the steps outlined above are described in more detail over the following subsections. Figure 2.21 shows an overview of how vertical columns (modelled and measured) along with filters and related data are created in this thesis. As described in this chapter, satellite and modelled data are combined to form recalculated corrected vertical columns (VCC), with masks created for potential anthropogenic, pyrogenic, and smearing influence (see Section 3.2.3)<sup>tb</sup>. The end product is a gridded data file that contains new and old satellite and modelled vertical columns, along with additional data to allow uncertainty calculation and other analyses. The original and recalculated AMF are also binned and stored.<sup>mh</sup>

## 2.6.2 Creating new shape factors

To visualise and analyse OMI HCHO columns, slant columns are transformed into vertical columns using the AMF. The shape factor ( $S$ ) is one of the key components in creation of the AMF (see Section 2.5, Equation 2.16). The shape factor is calculated using GEOS-Chem midday output (see Section 2.3.6.1) which provides simulated HCHO concentration profiles ( $\eta(z)$ ) and total columns ( $\Omega$ ) at  $2^\circ \times 2.5^\circ$  horizontal resolution. Using Equation 2.14 to determine the shape factor is straightforwards  $S(z) = \frac{\eta(z)}{\Omega}$ . The associated OMI per-pixel scattering weights are not changed in this calculation (unlike in Section 2.6.3.2).

Model output is provided as mixing ratios ( $C$  in ppbv), and is converted before being used in the shape factor calculation. The following equation converts model profile output from ppbv into number density ( $\eta$ ) in molecules  $\text{cm}^{-3}$ :

$$\eta_{HCHO} = C_{HCHO} \times \eta_a \times 10^{-9} \quad (2.21)$$

where  $\eta_{HCHO}$  is the number density of HCHO,  $\eta_a$  is the number density of air (from model output), and  $C_{HCHO}$  is the mixing ratio of HCHO. The modelled total vertical column  $\Omega_{HCHO}$  is determined by:

$$\Omega_{HCHO} = \Sigma_z (\eta_{HCHO} \times H(z))$$

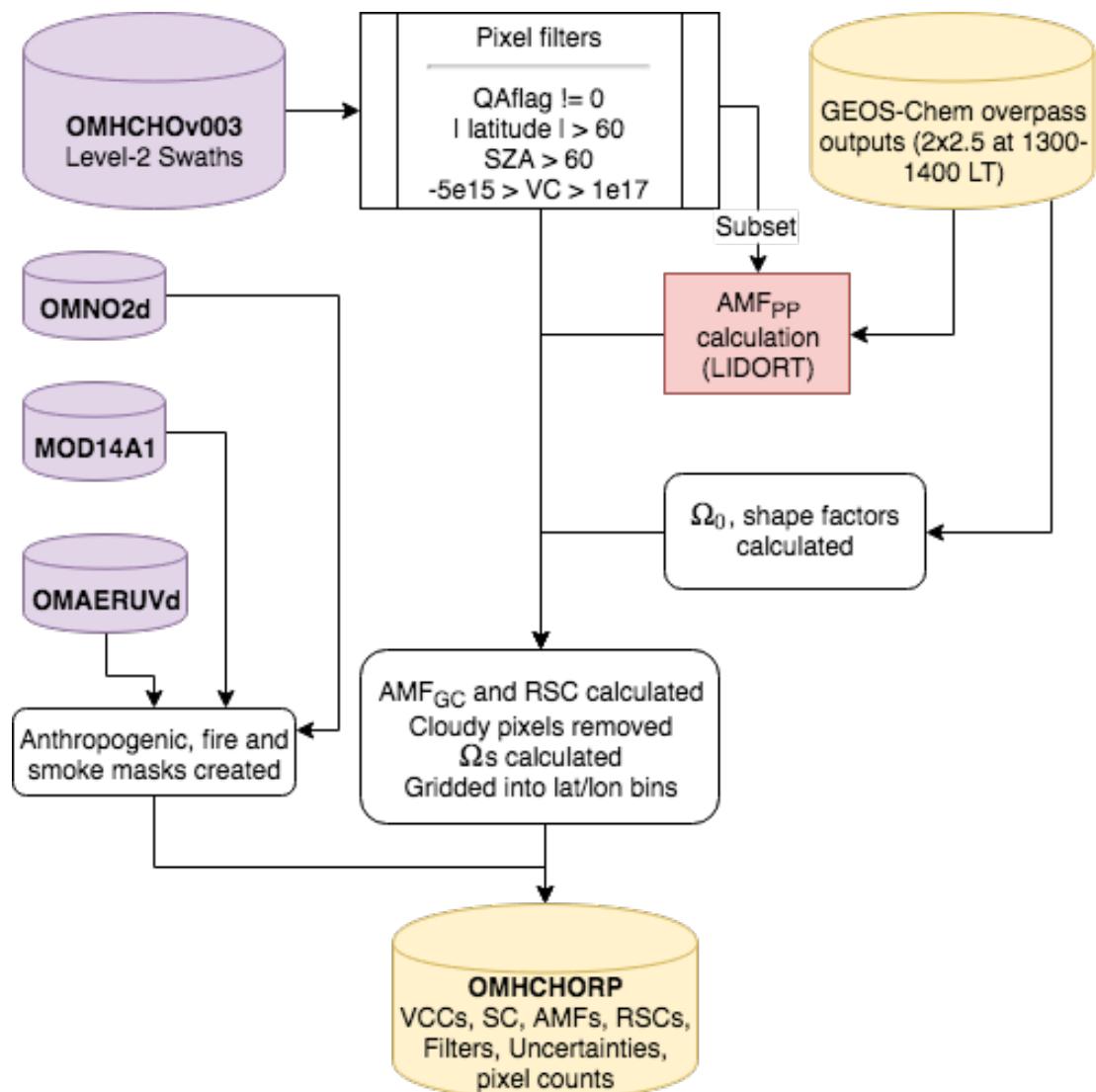


FIGURE 2.21: Flow diagram showing how OMHCHO level two swath data is read, processed, and gridded in this thesis.

where  $H(z)$  is the box height for altitude  $z$ . In effect this equation sums over the partial columns ( $\text{molec cm}^{-2}$ ) in each altitude level.

As a sanity check  $S_\sigma$  is calculated (through Equation 2.17) to confirm that these shape factors are equivalent. Comparing the resulting AMFs created by Equations 2.18 and 2.16 for each pixel provides confidence in the unit conversions (and other factors) applied. These AMF values are nearly identical in practice, with  $< 1\%$  bias and correlation coefficient  $r > 0.99$ .

### 2.6.3 Creating new AMF using GEOS-Chem

#### 2.6.3.1 Updating shape factors

From Equation 2.16 we have<sup>mh</sup>:

$$AMF = \int_0^{\infty} \omega(z) S(z) dz$$

Using the  $\omega(z)$  from satellite data, along with ~~our~~<sup>mh</sup> calculated  $S_z$  interpolated linearly onto the same vertical grid as  $\omega(z)$ , the AMF can be determined through integration. The integration is performed using a simple rectangular method, which multiplies the integrand midpoints by the change in height, and then takes the sum for each vertical box. This assumes that the provided scattering weights and shape factors are linear between the 47 resolved values.

#### 2.6.3.2 Recalculating the AMF

The major limitation of vertical columns that implement  $AMF_{GC}$  is that the scattering weights also depend on the model atmosphere, which is not updated when only the shape factor is recalculated. This is addressed using Fortran code written by Paul Palmer and Randal Martin, subsequently updated by Luke Surl (referred to here as PP code). This Fortran code is computationally expensive, and is run on a subset of the globe ( $50-10^\circ$  S,  $160^\circ$  W- $160^\circ$  E) covering Australia and the Pacific ocean. This allows both vertical column recalculation and reference sector correction.<sup>..</sup> The instrument sensitivity (or scattering weights;  $\omega$ ) and shape factors for each pixel are calculated within the PP code.

The code uses a combination of GEOS-Chem a priori profile information and satellite measurement data to calculate the  $AMF_{PP}$  by using LIDORT radiative transfer calculations to determine scattering.

Code for recalculating AMF using satellite swaths and modelled aerosol optical depths and gas profiles can be found at [http://fizz.phys.dal.ca/~atmos/martin/?page\\_id=129](http://fizz.phys.dal.ca/~atmos/martin/?page_id=129). The coded method is detailed in Palmer et al. (2001), with modifications for clouds and use of the LIDORT radiative transfer model (Spurr 2002) as described by Martin (2003). Modifications performed by Luke Surl at University of Edinburgh enabled the PP code to utilise OMI satellite data. Additionally, required tropopause heights averaged within satellite overpass times are created by modifying the GEOS-Chem diagnostic output. The calculation of  $\omega(z)$  is simplified by using large “look up tables” of values based on many parameters such as cloud top heights and optical depths. These AMF look up tables can be found in the source

code at <https://github.com/LukeSur1/amf581g>. The PP code uses HCHO concentration profiles averaged between 13:00 and 14:00 local time, including optical depths at 550 nm, and dust concentrations from GEOS-Chem, along with a subset of the OMI pixel information. The required information is taken from OMHCHO daily swath files, and sorted into csv files (one per day) which are then read by the PP code in conjunction with the GEOS-Chem outputs for each day. The PP code then produces a list of recalculated AMF values that are read by Python code and associated with the corresponding satellite pixel.

### 2.6.3.3 Saving the AMF with satellite pixels

OMI satellite pixels are read and filtered for quality (see Section 2.4.1) and have their AMFs calculated as shown above. This filtering removes highly uncertain pixels, including row anomaly affected pixels. Additional information is added to each pixel, including the new AMFs. Each pixel and its relevant data are saved in a long list for subsequent processing.

### 2.6.4 Vertical columns from AMF

All that remains for recalculating the total vertical column using `theourmh` new a priori shape factor is to apply the new AMF to the slant columns and grid them onto `aourmh` chosen resolution. Each satellite pixel at this stage has an associated SC along with three AMFs: the original ( $AMF_{OMI}$ ), one with recalculated shape factors ( $AMF_{GC}$ ), and one completely recalculated using PP code ( $AMF_{PP}$ ). These are used to create vertical columns ( $\Omega$ ) through Equation 2.8:  $\Omega = SC / AMF$ .

Figure 2.22 shows a comparison between the three estimates of satellite  $\Omega$  (each using a different AMF) over Australia for 2005. Fully recalculated  $AMF_{PP}$  show a distinct bi-modal distribution compared against the approximately Gaussian distributions of the  $AMF_{GC}$  and the original  $AMF_{OMI}$ . Additionally there is a steep decline in mean  $AMF_{PP}$  after February that is not seen in the other AMFs, and for the rest of the year the three AMFs follow the same pattern with  $AMF_{PP}$  around 10% lower on average. While the explanation for this is beyond the scope of this thesis, it does show that a complete recalculation of both the shape factor and the scattering weights makes a large difference in the AMF calculation when compared to just recalculating the shape factor. From here onward the  $AMF_{PP}$  is used in VC calculations and analysis, unless otherwise specified.

### 2.6.5 Reference sector correction

On a medium to long time scale, OMI degradation needs to be accounted for. To remove effects from the deterioration of the satellite instrument, measurements over the remote Pacific (`eurmh` reference sector) are combined with GEOS-Chem simulations to create a reference sector correction (RSC). In this thesis `we usemh` OMI HCHO columns `are usedmh` to estimate isoprene emissions, and correcting background HCHO does not affect this calculation. The correction should reduce bias caused by satellite degradation, without impacting isoprene emissions estimation. The RSC corrects for several

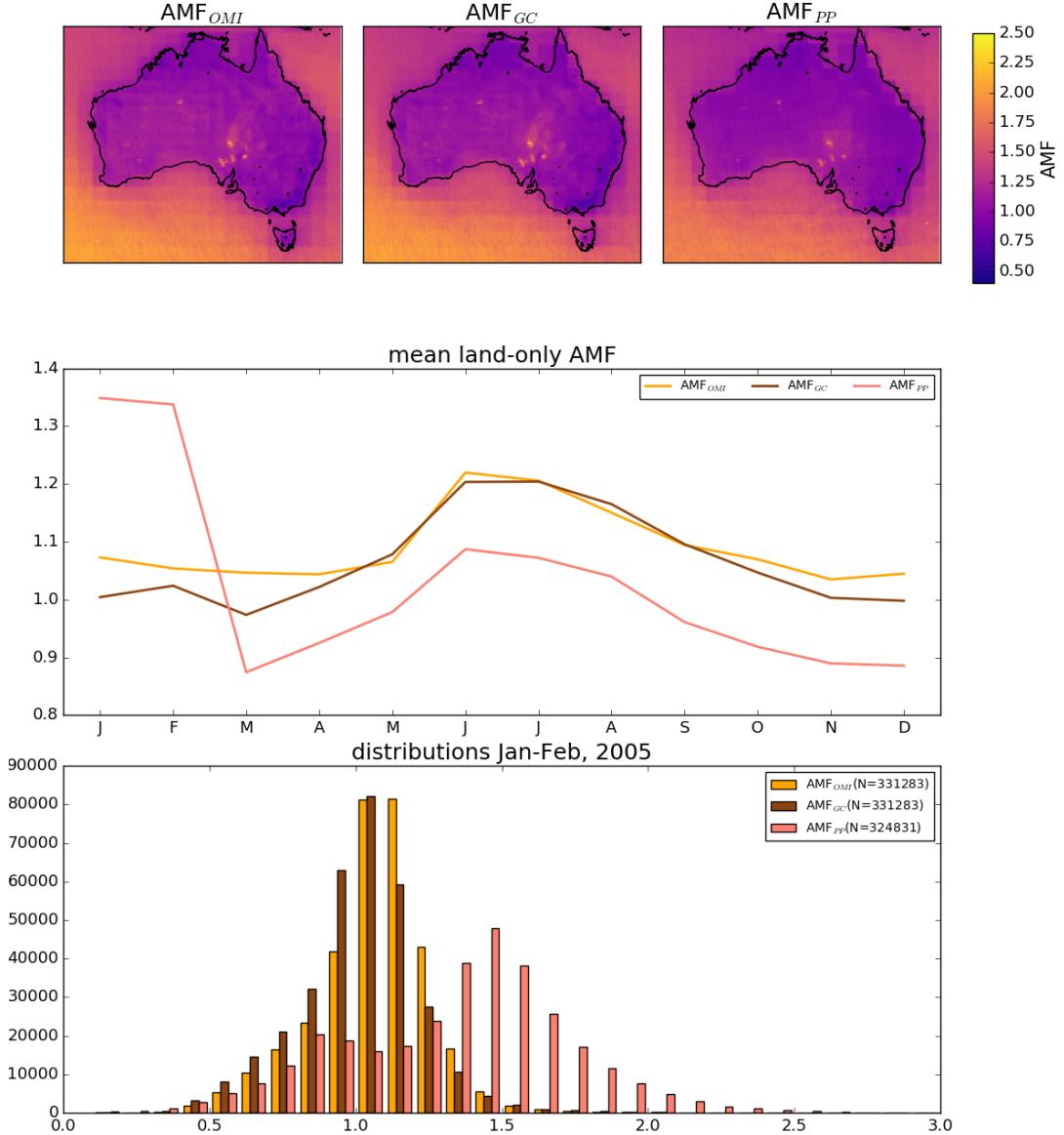


FIGURE 2.22: Top row: averaged OMI Satellite AMF for 2005, from the OMHCHO dataset (left,  $AMF_{OMI}$ ), recalculated using GEOS-Chem shape factors (middle,  $AMF_{GC}$ ), and recalculated using GEOS-Chem shape factors and scattering weights (right,  $AMF_{PP}$ ). Middle row: AMF time series over 2005 for each recalculation. Bottom row: AMF frequency distributions over January and February. Oceanic pixels are filtered out, only land pixels are included in this figure.

problems; however, it introduces some a priori model influence. One corrected problem is the potential influence of varying dead/hot pixel masks across the OMI 2-D detector array (De Smedt et al. 2015). This method also corrects for the errors introduced through correlations between BrO and HCHO absorption cross sections, which are especially significant at high latitudes (Gonzalez Abad et al. 2015).

HCHO products from OMI use a median daily remote Pacific Ocean radiance reference spectrum, over 15°S-15°N, 140°-160°W where it is assumed that the only significant source of HCHO is methane oxidation (De Smedt et al. 2008; Barkley et al. 2013; Kurosu and Chance 2014). Here a new reference sector correction is created using modelled and measured HCHO columns over the remote Pacific to produce corrected vertical columns (VCC). This follows Abad et al. (2016), and defines the remote Pacific as the band between 140°W to 160°W. Each satellite slant column measurement is corrected by how much the satellite reference sector measurements at that latitude diverge from modelled amounts over the reference sector.

Modelled slant columns over the reference sector ( $SC_{GC}$ ) are calculated by multiplying modelled vertical columns with the AMF calculated in prior sections using Equation 2.8:

$$SC_{GC} = \Omega_{GC} \times AMF$$

The longitudinal average is taken within the remote Pacific, as corrections are (assumed to be) longitudinally invariant. These modelled slant columns are averaged over the month and interpolated latitudinally to 500 equidistant bins. Figure 2.23 shows an example of the simulated reference sector vertical columns, calculated for January 1st 2005. In this figure the latitudinal resolution is increased from 2° to 0.36°, through linear interpolation, in order to form 500 vertical bins that are used in correcting the satellite data. This resolution is chosen to match that of Abad et al. (2016).

For OMI swaths, each row of measured data contains 60 “Across track” (track) measurements. The track index  $i$  relates the measurement to one of the 60 columns of data. Corrections (molecules cm<sup>-2</sup>) for each measurement are calculated by taking the difference between the measured slant column and the a priori slant column as follows:

$$Correction(i, j) = SC(i, j) - \Omega_{GC}(lat(j)) \times AMF(i, j) \quad (2.22)$$

where  $j$  represents a latitude index. The RSC is this list of *Correction* values binned by latitude using medians, and used per pixel based on the linear interpolation to that pixels latitude. The RSC is independently calculated for each of the 60 tracks, at each latitude in the 500 0.36° bins. This provides a different RSC for each of the three AMFs. Due to incomplete latitudinal coverage, the correction for each track is interpolated linearly between measurements, with corrections outside of the highest measured latitudes being equal to the corrections at the highest measured latitudes.

Figure 2.24 shows an example of the 60 track corrections for January 1st 2005. The points are satellite measurements longitudinally averaged over the remote Pacific, coloured by track number. Another way to look at this correction is given in the OMI corrections panel of Figure 2.23, which has tracks along the x axis and latitude on the y axis. This figure shows how corrections are distributed (over this 8-day sample) with more negative values towards the outside tracks, especially in the tropics.

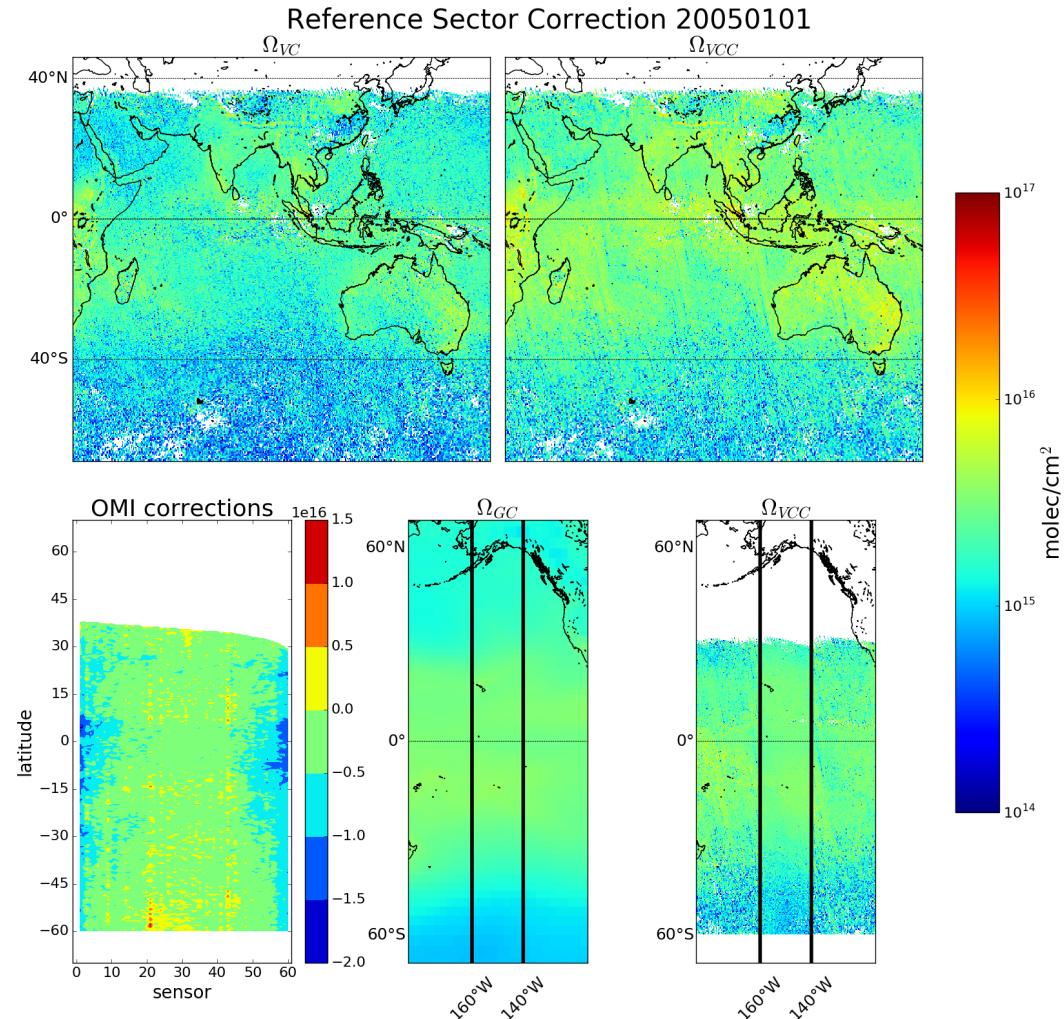


FIGURE 2.23: Example of remote Pacific RSC using 8-day average measurements and one month modelled data.  $\Omega_{VC}$  shows the uncorrected vertical columns, while  $\Omega_{VCC}$  shows the corrected vertical columns. OMI corrections shows the correction applied globally based on latitude and OMI track number (sensor).  $\Omega_{GC}$  shows the GEOS-Chem modelled HCHO VC over the reference sector (region within black vertical lines), with  $\Omega_{VCC}$  showing the corrected OMI VC over the same area.

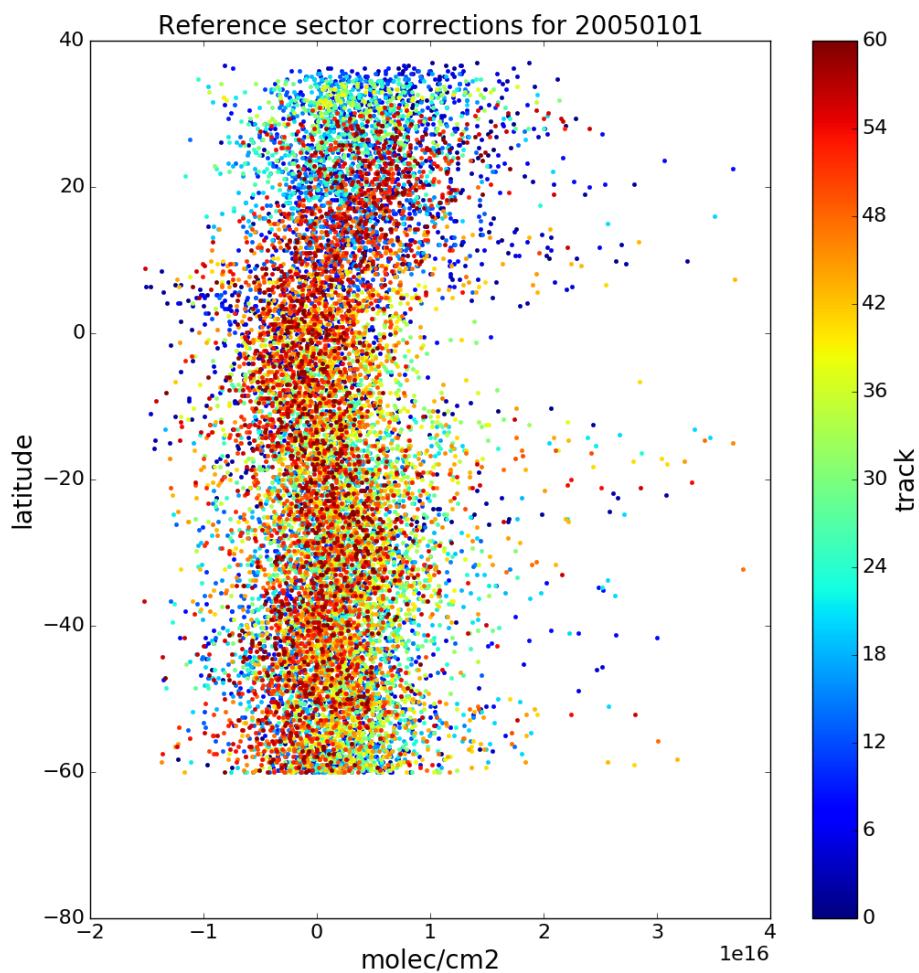


FIGURE 2.24: Example of track correction interpolation for January 1st 2005. Points represent the difference between satellite slant column measurements and modelled slant columns over the remote Pacific.

### 2.6.6 Corrected vertical columns

Corrected vertical columns, or  $VCC$ , are created using the difference between slant columns ( $SC$ ) and reference sector corrections ( $RSC$ , detailed in prior section) divided by the AMF.

$$VCC = \frac{(SC - RSC)}{AMF} \quad (2.23)$$

This is equivalent to taking the difference between the slant column and its measured reference sector equivalent ( $SC_0$ ), and then adding the modelled reference sector column ( $\Omega_{GC,0}$ ):

$$VCC = \frac{(SC - SC_0)}{AMF} + \Omega_{GC,0} \quad (2.24)$$

This method is used in several works, including De Smedt et al. (2008), De Smedt et al. (2012), De Smedt et al. (2015), Barkley et al. (2013), and Bauwens et al. (2016). Recently this correction was expanded (for OMI data) to include latitudinal and instrument track influence by Gonzalez Abad et al. (2015).

One *Correction* (Equation 2.22) is associated with every good satellite measurement. This is used to create a reference sector corrected measurement ( $VCC$ ) through the following equation:

$$VCC(i,j) = \frac{SC_{HCHO}(i,j) - correction(i, lat(j))}{AMF(i,j)} \quad (2.25)$$

For each good satellite measurement the corrected vertical column is calculated three times, once using each new AMF.

### 2.6.7 Binning the results daily

Finally the pixels are binned into a gridded dataset named OMHCHORP. The resolution is chosen to match the native resolution of the GEOS meteorological fields ( $0.25^\circ \times 0.3125^\circ$ ) to enable possible future comparison to high-resolution GEOS-Chem output (although this comparison is not performed in this thesis). A bin entry count is used to allow easy re-binning, and can be used to check for sparse data days due to filtering. Data averaged into this dataset include:

1. OMI slant column ( $SC$ )
2. OMI air mass factor ( $AMF_{OMI}$ )
3. OMI vertical column ( $VC_{OMI}$ )
4. OMI corrected vertical column ( $VCC_{OMI}$ )
5. GEOS-Chem recalculated air mass factor ( $AMF_{GC}$ )
6. GEOS-Chem recalculated vertical column ( $VC_{GC}$ )
7. GEOS-Chem recalculated corrected vertical column ( $VCC_{GC}$ )
8. GEOS-Chem air mass factor recalculated using PP code ( $AMF_{PP}$ )

9. GEOS-Chem corrected vertical column based on PP code ( $VCC_{PP}$ )
10. satellite pixel counts (summed into bins)
11. OMI vertical column fitting uncertainty

This process requires processing time and storage space, and is performed on the National Computational Infrastructure (NCI) supercomputer. In order to reprocess one year of swath files ( $\sim 162 \text{ GB} = 142 + 16 + 4 \text{ GB}$  OMHCHO, MOD14A1, and OMNO2d respectively per year), daily data were downloaded and then transformed into  $\sim 8 \text{ GB}$  (per year) of daily gridded data. This takes around 90 minutes per day; however, each day is completely independent and can be run in parallel once model output is available. Initially parallelism was built into the python code; however, simply running python code independently for each date was simpler and more scalable. As much as possible this work uses HDF-5 or NetCDF-4 formats, although GEOS-Chem output is in binary (bpch) format. The scripts to regrid and reprocess the swath dataset are available from github at [https://github.com/jibbals/OMI\\_regridding](https://github.com/jibbals/OMI_regridding).

### 2.6.8 Difference between original and corrected OMI HCHO columns

Corrected vertical columns ( $VCC$ ) of HCHO are created at  $0.25^\circ \times 0.3125^\circ$  horizontal resolution. Three  $VCC$  are created based on OMI and the new AMFs (from section 2.6.4) and RSCs. Figure 2.25 shows how the recalculated columns compare to the original (OMI, left) with January 2005 used as an example.  $VCC_{OMI}$  (original AMF) are the lowest overall for the depicted month, and recalculating just the shape factor increases the column amounts ( $VCC_{GC}$ ); however, recalculating both the shape factor and the scattering weights leads to a broader spread of values. Both sets of recalculated columns ( $VCC_{GC}$ ,  $VCC_{PP}$ ) show more high value ( $> 1.5 \text{ molec cm}^{-2}$ ) columns and lower background values over the ocean. This likely cause is the more recent modelled HCHO profiles used in the new calculations.

When reading  $AMF_{PP}$ , missing and near zero ( $AMF_{PP} < 0.0001$ ) values are removed, which leads to fewer overall pixels used to recalculate  $VCC_{PP}$  than either  $VCC_{OMI}$  or  $VCC_{GC}$ . Figure 2.26 shows how many pixels go into the column recalculation over Australia each month. The filtered pixel counts refer to how many pixels are removed through applied filters, which are described in the following section (Section 3.2.3). A decrease in pixel counts is apparent starting in April and lasting until August, which is caused by the reduction in good pixels recorded at increased solar zenith angles that occur during winter at higher latitudes. There is a consistent bias of 1 – 10% between the techniques, caused by the extra filtering applied to unreasonable AMFs calculated by the PP code.

Figure 2.27 shows global and Australian HCHO averaged total column maps for 2005, along with time series and distributions. The completely recalculated  $VCC_{PP}$  has a higher number of high HCHO amounts compared to both the original and partially recalculated  $VCC$ s. Both recalculated  $VCC$ s appear to have lower average background amounts, seen over the ocean and parts of southern Australia, and also increased average amounts in the HCHO hot spots around the north and eastern coastal regions. This comparison shows reprocessing with an updated model can have a systematic influence on the total column. Enough pixels remain to form monthly column amounts

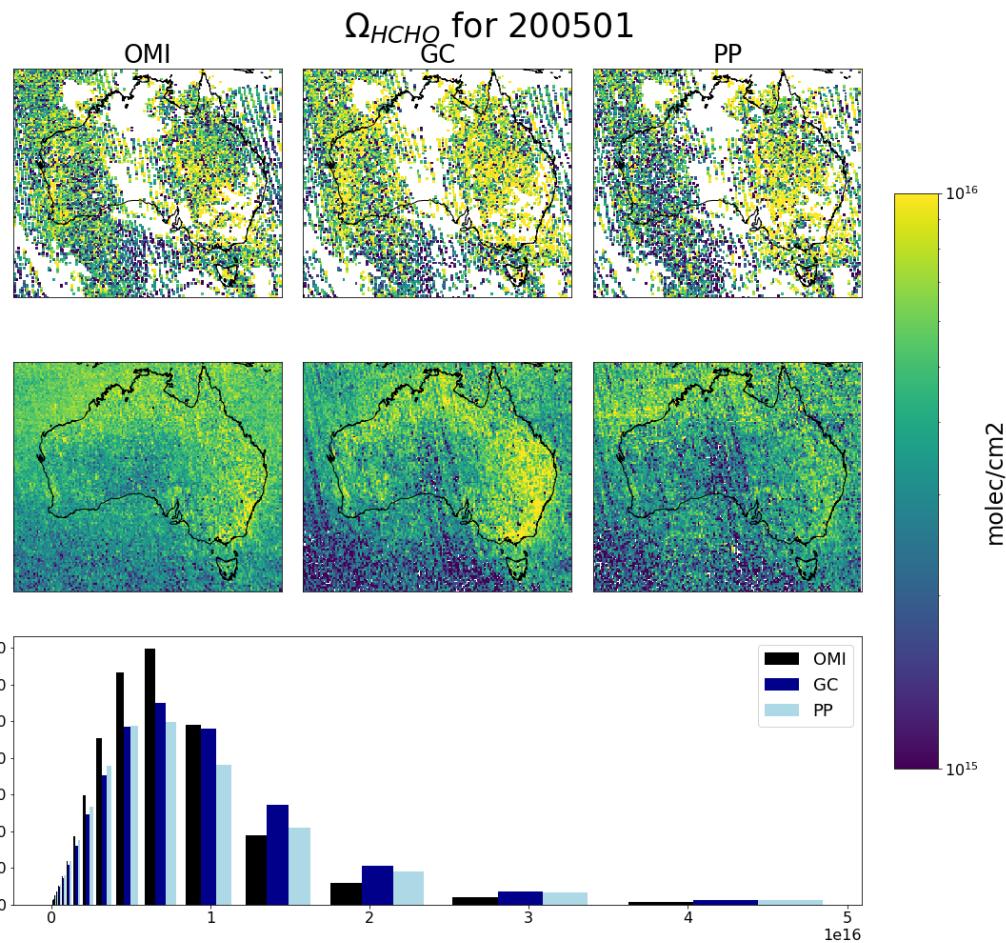


FIGURE 2.25: Row 1: regridded corrected  $\Omega_{HCHO}$  from OMHCHO on January 1, 2005: original (left), recalculated using new shape factors (middle), and additionally using updated scattering weights (right). Row 2: same but for the monthly average for January 2005. Row 3: the distribution over the month for each of the three column amounts. Distribution bins are logarithmic, resulting in wider bins for higher column amounts.

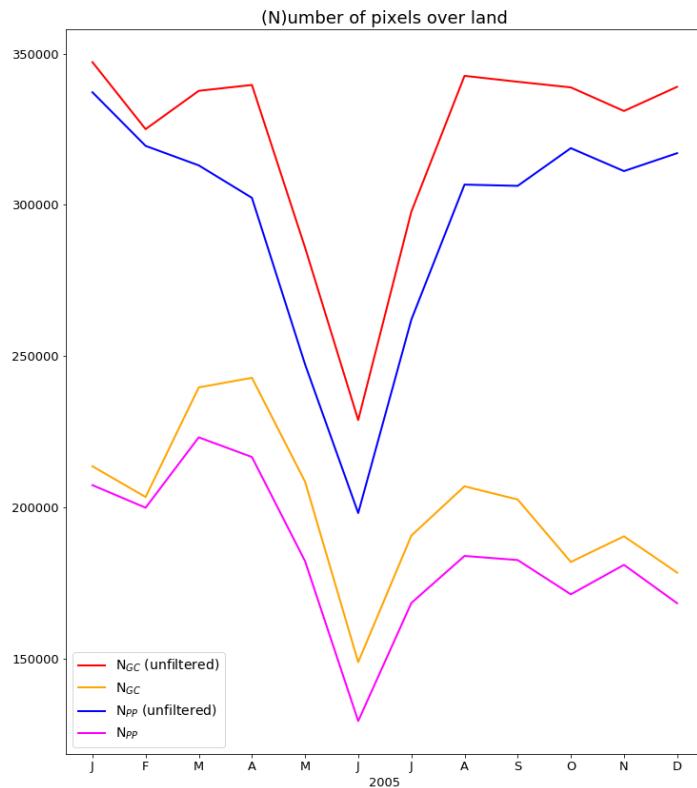


FIGURE 2.26: Monthly pixel counts over Australia (non-oceanic) used in recalculation of vertical columns.  $N_{GC}$  refers to the number of pixels used in AMF recalculation without running PP code,  $N_{PP}$  refers to those that have been recalculated using PP code. Filters applied are described in Section 3.2.3.

with good horizontal resolution, and fully recalculated, filtered columns ( $VCC_{PP}$ ) are used in the rest of this thesis unless otherwise specified.

~~Section 2.7 is moved into Chapter 3: Section 3.2.3<sup>tb</sup>~~

~~Section 2.8 is removed, text and figure moved to end of Section 2.6.1<sup>mh</sup>~~

## 2.7 Contributions and acknowledgements

All model runs, configuration, and alterations were performed by the Author, with initial discussions with Jenny Fisher. Satellite products were downloaded, collated, and (where appropriate) recalculated by the author. Recalculation of the OMHCHO data product is in part based on code written by Paul Palmer and Randal Martin, with updates and usage instructions given by Luke Surl. Output created in this chapter that is used outside of this thesis includes a filtered and regridded (but not recalculated) version of OMHCHO used in Kaitlyn J. Lieschke et al. (2019). “Decreasing trend in formaldehyde detected from 20-year record at Wollongong, Southeast Australia”. In: *Geophysical Research Letters*. DOI: 10.1029/2019GL083757. URL: <https://doi.org/10.1029/2019GL083757>, model run analyses used to debug GEOS-Chem satellite parameterisation, highly negative satellite pixels analysis that has been shared with the OMHCHO product developer, and products used in Chapter 3 may form part of a publication in preparation: Greenslade J. W., Fisher J. A. et al. Top-down emissions of isoprene over Australia.<sup>mh</sup>

## 2.8 Data access

Here I list<sup>mh</sup> the datasets used in this thesis are listed<sup>mh</sup>, along with where they are from and where they are used it is from and where it is used.

**AIRS** Atmospheric Infra-red Sounder instrument aboard the Aqua satellite (Texeira 2013). Satellite measurements of carbon monoxide downloaded from <https://search.earthdata.nasa.gov> with the product name AIRS3STD, DOI: 10.5067/AQUA/AIRS3STD. Data from this product are used in Chapter 4, and briefly described in Section 2.2.1.5.

**CPC** Climate Prediction Center global temperature data provided by the NOAA/OAR/ESRL PSD, Boulder, Colorado, USA, from their website at <https://www.esrl.noaa.gov/psd/>. Surface temperatures from this dataset are used in Section 2.6.8, and described in Section 2.2.2.3.

**ERA1** European Centre for Medium-range Weather Forecasts (ECMWF) Interim Re-analysis (ERA-I) (Dee et al. 2011). Downloaded using the online portal: <https://apps.ecmwf.int/datasets/data/interim-full-daily/>. These data are used in Chapter 4

**MUMBA** Measurements of Urban, Marine and Biogenic Air campaign (Paton-Walsh et al. 2017). Several trace gases from this campaign are examined here in Section 3.3.3, the dataset is described in more detail in Section 2.2.3.1.

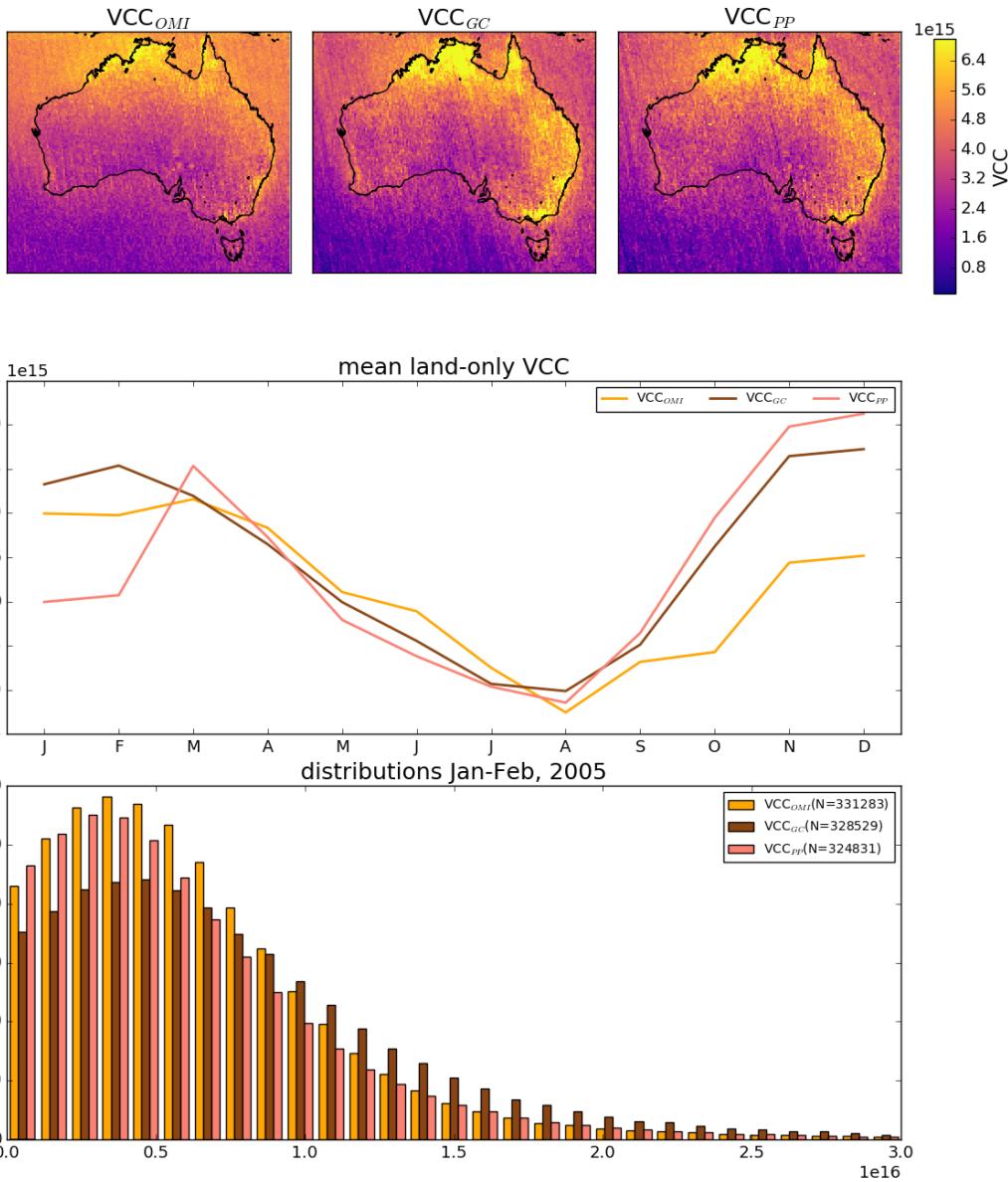


FIGURE 2.27: Top row: averaged OMI satellite VCC for 2005, from the OMHC dataset (left,  $VCC_{OMI}$ ), recalculated using GEOS-Chem shape factors (middle,  $VCC_{GC}$ ), and recalculated using GEOS-Chem shape factors and scattering weights (right,  $VCC_{PP}$ ). Middle row: VCC time series over 2005 for each recalculation. Bottom row: VCC frequency distributions over January and February. Oceanic pixels are filtered out, so only land pixels are included in this figure.

**OMHCHO** Satellite swaths of HCHO slant columns downloaded from <https://search.earthdata.nasa.gov>, with DOI 10.5067/Aura/OMI/DATA2015. More information can be found in Section 2.4.

**OMNO2d** Daily satellite NO<sub>2</sub> product downloaded from <https://search.earthdata.nasa.gov/search>, DOI:10.5067/Aura/OMI/DATA3007. For more information in refer to section 2.2.1.2.

**OMAERUVd** Gridded satellite based AAOD measurements downloaded from the NASA earth data portal <https://search.earthdata.nasa.gov>. A summary can be found at [https://disc.gsfc.nasa.gov/datasets/OMAERUVd\\_V003/summary](https://disc.gsfc.nasa.gov/datasets/OMAERUVd_V003/summary), Omar O. Torres (2008). *OMI/Aura Near UV Aerosol Optical Depth and Single Scattering Albedo L3*. DOI: 10.5067/Aura/OMI/DATA3003. URL: [https://disc.gsfc.nasa.gov/datasets/OMAERUVd\\\_\\\_V003/summary](https://disc.gsfc.nasa.gov/datasets/OMAERUVd\_\_V003/summary) (visited on 05/30/2019). This dataset is described in detail in Section 2.2.1.3.

**SPS** Observations from Westmead Air Quality Station, stage 1 (SPS1, Keywood et al. (2016a)) from 5 February to 7 March 2011, stage 2 (SPS2, Keywood et al. (2016b)) from 16 April to 14 May 2012. SPS1 is available from <https://doi.org/10.4225/08/57903B83D6A5D>, and SPS2 from <https://doi.org/10.4225/08/5791B5528BD63>. Some trace gas measurements are examined here in Section 3.3.3. Relevant measurements described in detail in Section 2.2.3.1.

**Wollongong FTIR** The instrument is part of the Network for the Detection of Atmospheric Composition Change (NDACC) and data can be retrieved from the NDACC database (<http://www.ndaccdemo.org/stations/wollongong-australia>). The current principle investigator producing and quality assuring the dataset is Dr. Nicholas Jones.

**Ozone sondes** Ozonesonde data were retrieved from the World Ozone and Ultraviolet Data Centre (WOUDC) <http://woudc.org/data/explore.php>.



## Chapter 3

# Biogenic Isoprene Emissions in Australia

### 3.1 Introduction

Biogenic volatile organic compounds (BVOC) affect the oxidative capacity of the atmosphere and their emissions are largely driven by what type of vegetation is in the area (Kefauver, Filella, and Peñuelas 2014). In the troposphere, BVOC emissions affect hydroxyl radical ( $\text{OH}$ ) cycling, ozone ( $\text{O}_3$ ) production, secondary organic aerosol (SOA) production, and methane lifetime. Australian forests are strong emitters of isoprene, the primary BVOC emitted globally (Guenther et al. 2006; Messina et al. 2016). Isoprene is relatively difficult to measure due to its high reactivity and short lifetime. Poor measurement coverage of isoprene, isoprene products, and isoprene emissions within Australia means that emissions are poorly understood. The lack of knowledge about emissions makes it difficult to estimate the subsequent atmospheric processes.

Emission models used to derive estimates of isoprene fluxes are based on understanding the emissions from different plant species (phenotypes) in varying conditions (see Section 1.3.2)<sup>tb</sup>. Guenther et al. (2012) estimated global biogenic isoprene emissions at roughly  $535 \text{ Tg yr}^{-1}$ , while Sindelarova et al. (2014) estimated around  $411 \text{ Tg yr}^{-1}$ . Reactions following emissions are complex, and are sensitive to other trace gases in the atmosphere. Uncertainties in several important products such as ozone and SOA are increased due to both isoprene measurement difficulties and its complicated subsequent chemical mechanisms. Isoprene emissions in the frequently used MEGAN model may be overestimated in Australia (see Section 1.3.1)<sup>tb</sup> since they are based on measurements taken from a few young trees (Winters et al. 2009) that may emit more than older trees (Emmerson et al. 2016). The sample of trees used to construct the MEGAN emissions model included 4 types of *Eucalyptus*, which are not representative of the hundreds of species that make up Australian forests. Additionally, how these species react to biological and meteorological stresses is unclear (Winters et al. 2009; Fortems-Cheiney et al. 2012). Emissions estimates are necessary inputs for atmospheric chemistry models and improving these estimates for Australia is a primary goal of this thesis.

This chapter describes and implements In this chapter, I describe and implement<sup>mh</sup> a *top-down* technique using satellite measurements of HCHO to calculate surface isoprene emissions. HCHO is a primary product of most BVOC (including isoprene) oxidation, and is measured by satellites via remote sensing. In situ isoprene concentration

measurements are costly and sparse within Australia, while satellite HCHO data are plentiful and freely available, making this technique very attractive. Top-down techniques have informed isoprene emission inventories in North America (Abbot 2003; Palmer et al. 2003; Palmer et al. 2006; Millet et al. 2006; Millet et al. 2008), South America (Barkley et al. 2013), Europe (Dufour et al. 2008; Curci et al. 2010), Africa (Marais et al. 2012), Asia (Fu et al. 2007; Stavrakou et al. 2014), India (Surl, Palmer, and Abad 2018), and even globally (Shim et al. 2005; Fortems-Cheiney et al. 2012; Bauwens et al. 2016). In this thesis the technique is applied~~I apply the technique~~<sup>mh</sup> focusing solely on Australia for the first time.

### 3.1.1 Aims

Recent work suggests that modelled emissions may be overestimated in southeast Australia (Emmerson et al. 2016). This chapter aims to improve the understanding of isoprene emissions over the whole of Australia, clarifying the spatial distribution of bias and how these biases impact modelled chemistry. Isoprene emissions are estimated~~I estimate isoprene emissions~~<sup>mh</sup> in Australia using a top-down technique based on OMI HCHO measurements and GEOS-Chem modelled yields. This a posteriori top-down estimate is evaluated against bottom-up a priori estimates and also briefly compared against available ground-based measurements. The GEOS-Chem model is modified to run with the a posteriori isoprene emissions to determine potential impact on modelled chemistry. Goodness of fit between in situ, satellite, and modelled HCHO is determined before and after scaling emissions estimates.

First an outline is provided showing~~I outline~~<sup>mh</sup> why current isoprene emissions estimates are inadequate and how they can be improved. This thesis discusses~~I discuss~~<sup>mh</sup> literature that shows how the estimates may be too high, and describes~~s~~<sup>mh</sup> how emissions may be calculated using satellite datasets. Section 3.2 lays out how new isoprene emissions are estimated, with results examined in Section 3.3. Section 3.3 includes a comparison of updated satellite HCHO columns (Chapter 2) to available measurements, and an examination of how these changes in emissions would affect ozone concentrations in Australia. Uncertainties for each step along the way are quantified in Section 3.4.

### 3.1.2 Existing emissions estimates

MEGAN is one of the most widely used sources for estimating biogenic isoprene emissions. However, along with other models that rely on measured plant emission rates, it is poorly calibrated for Australian conditions. Emissions of isoprene ( $C_5H_8$ ) appear to be overestimated within Australia (Sindelarova et al. 2014; Stavrakou et al. 2014; Bauwens et al. 2016; Emmerson et al. 2016), although the lack of measurements of isoprene emission rates in Australia makes this overestimation difficult to characterise. The bias may be as high as a factor of 6, and is not easy to fix (see Section 1.3.2). Bauwens et al. (2016) showed that isoprene emissions were overestimated by up to a factor of 3 over Australia. Emmerson et al. (2016) suggest isoprene emission estimates are 2-6 times too high compared against available measurements of isoprene concentrations. They compared modelled data to campaign measurements from multiple

sites over different seasons and found that scaling emissions did not universally improve model outputs globally.<sup>tb</sup>

Recently Bauwens et al. (2016) estimated isoprene emissions with a top-down technique using the IMAGESv2 global CTM. They calculated emissions that create the closest match between model and OMI satellite vertical columns, and compared these a posteriori data with their a priori (model) data and independent datasets. For Australia they found MEGAN based estimates ranging from 38 Tg C yr<sup>-1</sup>- 94 Tg C yr<sup>-1</sup>, and a posteriori emissions of 36 Tg C yr<sup>-1</sup>. This wide range demonstrates the effect of unknowns and uncertainties involved in isoprene emission estimation. Their results are compared to results here in Section<sup>tb</sup> 3.3.1. In this thesis I develop and analyse<sup>mh</sup> a top-down emission estimate is developed, analysed and, which is<sup>mh</sup> then compared against MEGAN. HI also look at how changed emissions affect modelled ozone levels over Australia is also looked into<sup>mh</sup>.

### 3.1.3 Top-down isoprene emissions estimates

In the remote troposphere HCHO production is dominated by methane oxidation, while in the continental boundary layer production is largely due to non-methane VOCs (NMVOCs) (Abbot 2003; Kefauver, Filella, and Peñuelas 2014). This leads to a causal relationship between enhanced HCHO concentrations and NMVOC emissions at low (< 1 km) altitudes. NMVOCs are generally short lived (< 1 hr), and the most prominent of these is isoprene. Isoprene is emitted and enters the atmosphere in the gas phase, where it begins a complex series of reactions. HCHO is produced with high yield in many reactions beginning with isoprene oxidation (discussed in more detail in Section 1.3.4) and it has a lifetime of a few hours (Kefauver, Filella, and Peñuelas 2014).

Top-down estimates determine emissions of a particular species through careful analysis of the measurable products of that species. This generally takes advantage of longer-lived products that may reach an equilibrium in the atmosphere. Continental tropospheric HCHO enhancements can be directly linked to biogenic isoprene emissions. Since 1997, when the Global Ozone Monitoring Experiment (GOME) satellite measurements were first used to measure HCHO, satellites have been used to provide a total column measurement of HCHO, enabling isoprene emissions estimation by top-down methods (Thomas et al. 1998; Palmer et al. 2001; Bauwens et al. 2016). Using satellite information to improve estimates of biogenic emissions has been highlighted as a valuable use of satellite derived datasets (Streets et al. 2013). Here NASA's OMHCHO product based on measurements from the OMI instrument onboard the Aura satellite (see Section 2.4) is the basis for a top-down biogenic isoprene emission estimate over Australia.

There are two top-down isoprene emission estimation techniques, Bayesian and linear, which are discussed briefly here. Both the linear and Bayesian techniques assume that modelled chemistry is accurate and only try to correct precursor emissions. This is an additional source of uncertainty given existing uncertainties in chemical mechanisms.

### 3.1.3.1 Bayesian inversion

Bayesian inversion optimises model parameters in order to minimise the difference between model output and an (ideally) independent dataset such as satellite measurements. Emissions of isoprene (and other precursors to HCHO) will form part of the set of model parameters that are adjusted to make the model HCHO output most closely match satellite measurements. These inversions can be set up to account for effects from transport and allow source attribution (e.g., Curci et al. 2010; Fortems-Cheiney et al. 2012).

In general, a model (the forward model) is used to determine the relationship between HCHO ( $y$ ) and the state variable  $x$ , which represents isoprene emissions (and other variable parameters of interest):

$$y = \mathbf{K}x + b + \epsilon \quad (3.1)$$

where  $\epsilon$  are the (assumed) independent errors in measurements.  $K$  is the Jacobian matrix determined from the forward model representing the sensitivity of  $y$  to the state variable  $x$ . Essentially the  $K$  matrix is the modelled estimation of how  $y$  responds to each of the driving parameters represented by elements of  $x$ . This  $K$  matrix is used in conjunction with error covariance in  $x$  to determine the most likely solution to  $x$ , given what is known about  $y$ .

This method was used by Shim et al. (2005) to optimise isoprene emissions in areas with high HCHO concentrations. They showed model underestimation of isoprene emissions by 14-46%, which was reduced to 3-25% after applying satellite based improvements. More recently Kaiser et al. (2018) showed a 40% bias in MEGAN isoprene emissions over the southeast US using a Bayesian inversion based on OMI HCHO.

An advantage (over the linear method described below) of the Bayesian method is that it can account for pyrogenic and anthropogenic emissions, as these form part of the state variable  $x$  as well as transport. However, biases in the underlying model will still propagate through to the a posteriori emission estimation (Curci et al. 2010). More limiting is the fact that the Bayesian method is computationally expensive. In this work [the Bayesian method is not used](#)<sup>I do not use the Bayesian method</sup> due to the computational costs surpassing the resources available.

### 3.1.3.2 Linear inversion

The linear technique is performed in this thesis. Vertical columns of HCHO from satellite and modelled yield from isoprene allow the inference of local (grid space) isoprene emissions (Palmer et al. 2003; Millet et al. 2006; Marais et al. 2012; Surl, Palmer, and Abad 2018). The primary assumption of the linear inversion technique is that HCHO and its precursors (primarily isoprene) are in a linear steady state relationship. This allows one to link isoprene emissions to HCHO measurements using production and loss rates. Essentially a linear relationship between total column HCHO ( $\Omega$ ) enhancement above a background level ( $\Omega_0$ ) and isoprene emissions ( $E_{isop}$ ) is determined:

$$\Omega = S \times E_{isop} + \Omega_0$$

This uses modelled vertical columns and emissions to estimate the slope ( $S$ ). Then this modelled  $S$  is applied to satellite measurements of  $\Omega$  ( $\Omega_{sat}$  and  $\Omega_{sat,0}$ ) to determine  $\hat{E}_{isop}$ :

$$\hat{E}_{isop} = \frac{\Omega_{sat} - \Omega_{sat,0}}{S}$$

This is described further in Section 3.2, with an outline in Section 3.2.1.

The calculation requires reaction rates and yields from isoprene to HCHO, which can be determined most readily using chemical modelling. The method for calculating isoprene emissions from HCHO is laid out in Palmer et al. (2003), taking into account the expected lifetime and reaction rates of the precursor VOCs and HCHO. In their work, isoprene emissions fluxes over the US were derived using the GOME satellite instrument. The method has since been applied to many regions using GOME, SCIAMACHY, OMI, and GOME-2 satellite data (e.g., Abbot 2003; Barkley et al. 2013; Stavrakou et al. 2014; Surl, Palmer, and Abad 2018).

The linear inversion assumes fast HCHO yield from isoprene and no precursor transport, which is unrealistic in certain scenarios. For example, high wind speeds can transport precursors, or low  $\text{NO}_x$  concentrations can slow HCHO production (Palmer et al. 2006; Surl, Palmer, and Abad 2018). Filtering out data that do not match assumptions is required but can limit the utility of this technique, and leads to some dependence on environmental factors. Uncertainties in the technique are discussed in more detail in Section 3.4.1. Nonetheless, a major benefit is that the simple nature of the inversion requires very little computational power after acquiring satellite and model datasets, even over large amounts of gridded data. This allows an inversion using more than 8 years of satellite and model data, capturing inter-annual variability over all of Australia. With the computational resources available this would not have been possible using the Bayesian inversion.

## 3.2 Methods

~~The broadly followed~~<sup>mh</sup>he method of Palmer et al. (2001) is broadly followed<sup>mh</sup> to create a biogenic isoprene emissions estimate over Australia. A relationship is modelled between biogenic-only midday tropospheric columns of HCHO and GEOS-Chem mid-day biogenic isoprene emission rates, and then this relationship is applied to satellite measured HCHO total columns to derive a new isoprene emissions estimate. Daily modelled values averaged between 13:00-14:00 LT are used to match the overpass time of the Aura satellite. Then the slope is calculated using reduced major axis regression between the a priori isoprene emissions (those from GEOS-Chem,  $E_{GC}$ ) and tropospheric HCHO columns in each model grid box each month. There is very little HCHO above the tropopause, so differences between total and tropospheric column are negligible. In this work total and tropospheric column HCHO are used interchangeably and referred to using  $\Omega$ .

### 3.2.1 Outline

This section provides an overview of the steps involved in creating a top-down emissions estimate. This process is summarised in Figure 3.1.

1. Corrected vertical columns ( $\Omega_{OMI}$ ; saved in the OMHCHORP dataset) are calculated (see Section 2.6) using level two OMI HCHO satellite data (see Section 2.4), along with GEOS-Chem model runs (see Section 2.3.6). Satellite columns are binned into both  $0.25^\circ \times 0.3125^\circ$  and  $2^\circ \times 2.5^\circ$  horizontal resolutions. In this step model background values (columns over the remote Pacific) are used to correct the vertical columns, which is explained in Section 2.6.5.
2. Level three satellite data are used to make anthropogenic, fire, and smoke influence masks (see Section 3.2.3). These are applied to remove  $\Omega_{OMI}$  that may be influenced by pyrogenic or anthropogenic sources.
3. A mask is created showing where the HCHO production is not dominated by local isoprene emissions. This is determined by calculating smearing over Australia using two model runs with differing isoprene emissions. The smearing value is determined as  $\hat{S} = \Delta\Omega_{GC}/\Delta E_{OMI}$ : the ratio of the differences between model runs of HCHO columns and isoprene emissions. A full description of the creation of this smearing filter is given in Section 3.2.3.3.
4. GEOS-Chem modelled biogenic emissions of isoprene ( $E_{GC}$ ) along with biogenic columns of HCHO ( $\Omega_{GC}$ ), both averaged over  $2^\circ \times 2.5^\circ$  horizontally and 13:00-14:00 LT temporally, are used to calculate a reduced major axis linear regression slope  $S$  ( $\Omega_{GC} = S \times E_{GC} + \Omega_{GC,0}$ ). Calculation of this modelled slope is explained in Section 3.2.5.
5. Satellite HCHO  $\Omega_{OMI}$  and  $S$  then form the basis for the top-down estimate of biogenic isoprene emissions ( $E_{OMI}$  atoms C  $\text{cm}^{-2} \text{ s}^{-1}$ ). This [is the product is our<sup>mh</sup>](#) a posteriori, and calculation details are given in Section 3.2.8.
6. A posteriori top-down emissions  $E_{OMI}$  are compared against a priori emissions, and analysed in conjunction with independent observations from in situ measurements (MUMBA and SPS). Results are examined in Section 3.3.
7. GEOS-Chem is run using the a posteriori emissions (see Section 3.2.9), and HCHO, O<sub>3</sub>, isoprene, and NO<sub>x</sub> outputs are compared to a priori values and to campaign and satellite measurements where possible (Sections 3.3.3).

### 3.2.2 Masks and reprocessed satellite HCHO

Satellite data pixels are read from OMHCHO, the level 2 OMI HCHO dataset, AMFs are recalculated, and then pixels are gridded daily into  $0.25^\circ \times 0.3125^\circ$  horizontal bins. This forms the intermediate product OMHCHORP, which is fully described in Section 2.6.1. This dataset includes gridded satellite HCHO columns ( $\Omega_{OMI}$ ), along with pixel counts (how many satellite data points were used for each grid box) to allow averaging, re-binning, and uncertainty analysis. In this thesis [the OMI product is used to use the OMI product<sup>mh</sup>](#) as the overpass time (13:30 local time) is closer to peak isoprene emissions than that of GOME or GOME-2.

In order to determine biogenic HCHO enhancements from  $\Omega_{OMI}$ , filters for non-biogenic sources are required. These masks are described in Section 3.2.3, and a brief

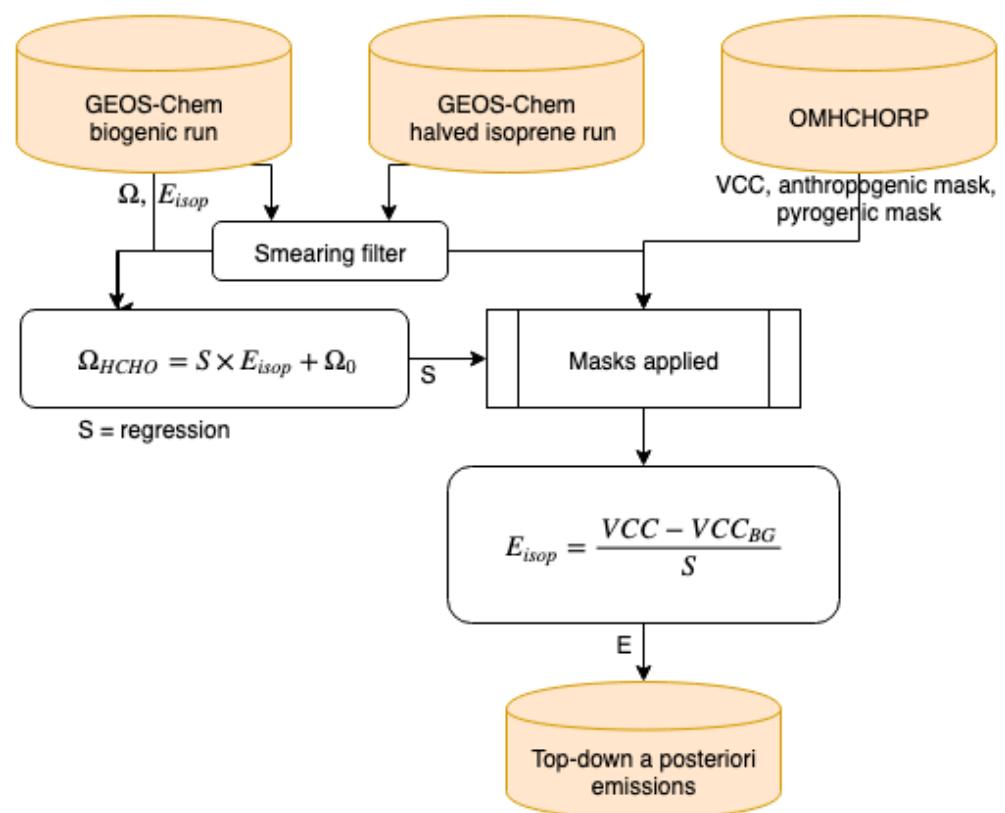


FIGURE 3.1: Top-down isoprene emissions estimate formation using OMHCHORP and GEOS-Chem outputs.

[overviewrecap<sup>tb</sup>](#) is provided here. Methane oxidation is a major part of the formation of background HCHO; however, the linear regression used to estimate isoprene emissions effectively removes this source (by assuming a constant background), which means a methane contribution filter is not required. Anthropogenic, pyrogenic, and smoke influence masks are created from three satellite products: NO<sub>2</sub> from OMNO2d, fire counts from MOD14A1, and AAOD from OMAERUVd, respectively.

1. The fire mask is created daily using non-zero MODIS fire counts over the prior 2 days that occur in local or adjacent grid squares at  $0.25^\circ \times 0.3125^\circ$  horizontal resolution.
2. Influence from transported smoke plumes is removed by flagging grid squares where OMI aerosol absorption optical depth (AAOD, from OMAERUVd) is greater than 0.03.
3. A filter for anthropogenic influence is created daily using OMNO2d NO<sub>2</sub> tropospheric column amounts, masking any grid squares with greater than  $2.0 \times 10^{15}$  molec cm<sup>-2</sup> on any particular day, along with grid squares where the yearly average is above  $1.5 \times 10^{15}$  molec cm<sup>-2</sup>.

The recalculated corrected vertical columns are saved to OMHCHORP dataset both before and after applying the filters to allow filter analysis.

### 3.2.3 Filtering Data

[This subsection \(3.2.3\) has been moved here from Chapter 2. I have added this to be removed paragraph instead of highlighting everything.<sup>tb</sup>](#)

A major goal in this work is to infer biogenic isoprene emissions from HCHO columns. Isoprene is the dominant source of HCHO over continental land masses; however, other precursor NMVOCs can contribute to observed HCHO. The main interference in the biogenic isoprene signal in HCHO comes from fire smoke plumes and major anthropogenic source regions. Biomass burning can be a large local or transported (via smoke plumes) source of HCHO, glyoxal (CHOCHO), and other compounds that influence HCHO levels. Anthropogenic emissions from power generation, transport, and agriculture can influence these levels as well. So to infer biogenic isoprene emissions, pyrogenic and anthropogenic influences should be removed (where possible) from modelled and measured data. In GEOS-Chem in this thesis, pyrogenic and anthropogenic emissions are turned off in the model run used to estimate biogenic yields. In the OMI HCHO satellite product, potentially pyrogenic or anthropogenic affected pixels are masked.

In this work anthropogenic and pyrogenic influences on the OMHCHO satellite HCHO columns are removed by masking active fires, high AAOD, and high NO<sub>2</sub> levels measured by satellite. Active fires and suspected smoke plumes are masked, and together termed the pyrogenic filter. NO<sub>2</sub> measurements are used to mask potential anthropogenic influence. Details of each filter are provided in the following sub-sections. A summary of yearly filtering over Australian land squares at  $0.25^\circ \times 0.3125^\circ$  resolution is provided in Table 3.1. Figure 3.2 shows an example year of anthropogenic and pyrogenic filtering, highlighting how many days and pixels are removed across Australia. The anthropogenic filter completely removes grid squares

TABLE 3.1: Satellite pixels remaining after filtering by active fires, smoke, and anthropogenic masking. In parenthesis are the portion of pixels filtered.

<b>Year</b>	<b>Pixels<sup>a</sup></b>	<b>Active fires</b>	<b>Anthropogenic</b>	<b>Smoke</b>	<b>Total<sup>b</sup></b>
2005	3.9e+06	3.5e+06( 9.2%)	3.8e+06( 1.3%)	3.8e+06( 1.7%)	3.4e+06(11.5%)
2006	3.8e+06	3.4e+06(11.2%)	3.7e+06( 2.4%)	3.7e+06( 3.2%)	3.2e+06(14.8%)
2007	3.7e+06	3.4e+06( 9.9%)	3.7e+06( 2.0%)	3.7e+06( 2.6%)	3.3e+06(13.0%)
2008	3.5e+06	3.3e+06( 8.1%)	3.5e+06( 1.4%)	3.5e+06( 1.1%)	3.2e+06(10.0%)
2009	2.7e+06	2.4e+06( 9.1%)	2.6e+06( 1.8%)	2.6e+06( 1.7%)	2.4e+06(11.6%)
2010	2.0e+06	1.9e+06( 6.5%)	2.0e+06( 1.4%)	2.0e+06( 0.5%)	1.9e+06( 8.1%)
2011	1.9e+06	1.7e+06(12.5%)	1.9e+06( 1.9%)	1.9e+06( 2.7%)	1.6e+06(15.2%)
2012	2.5e+06	2.2e+06(14.3%)	2.5e+06( 1.8%)	2.5e+06( 3.3%)	2.1e+06(17.4%)

a: how many land pixels are read over Australia, after cloud fraction filtering.

b: Total removed pixels accounts for overlap between filters.

over Brisbane, Melbourne, and Sydney. Other major cities in Australia either do not emit enough NO<sub>2</sub> to be clearly identifiable by satellite or are too dispersed and do not breach the threshold to be filtered as anthropogenic. The anthropogenic filter removes from  $\sim 0.25\% - \sim 5\%$  of grid squares each day. Pyrogenic filtering removes from  $\sim 5\% - \sim 20\%$  of the available grid square measurements per day. This filter tends to cover forested areas (as they are more prone to burning) as well as some hot spots that are likely due to gas flaring or burning.

### 3.2.3.1 Pyrogenic filter

MODIS fire counts are used in conjunction with smoke AAOD enhancements from OMI to remove data points that may be affected by fires or fire smoke plumes. The MODIS fire counts come from a combination of measurements from the Terra and Aqua satellites (Terra overpasses at 10:30, 22:30; Aqua at 13:30, 01:30 local time). Satellite AAOD from product OMAERUVd (described in Section 2.2.1.3) is analysed over Australia to determine a suitable filter threshold. AAOD is used instead of an alternative product AOD as it is less sensitive to the presence of clouds (Ahn, Torres, and Bhartia 2008).

OMHCHO total column HCHO  $\Omega$  is processed into a  $0.25^\circ \times 0.3125^\circ$  horizontal daily grid as described in previous sections. Pyrogenic filters are interpolated to the same horizontal resolution as  $\Omega$  to simplify application. The following steps are performed to create the pyrogenic influence mask:

1. MOD14A1 daily gridded Aqua/Terra combined fire counts ( $1 \times 1 \text{ km}^2$ ) are binned into  $0.25^\circ \times 0.3125^\circ$  bins (matching the resolution of binned  $\Omega$ ).
2. A rolling mask is formed that removes  $\Omega$  if one or more fires are detected in a grid square, or in the adjacent grid square, up to 2 days previously. This includes the current day, making 3 days of fires in total being filtered out on each day.
3. AAOD at 500 nm is mapped from OMAERUVd ( $1 \times 1^\circ$  resolution) onto the  $0.25^\circ \times 0.3125^\circ$  resolution.

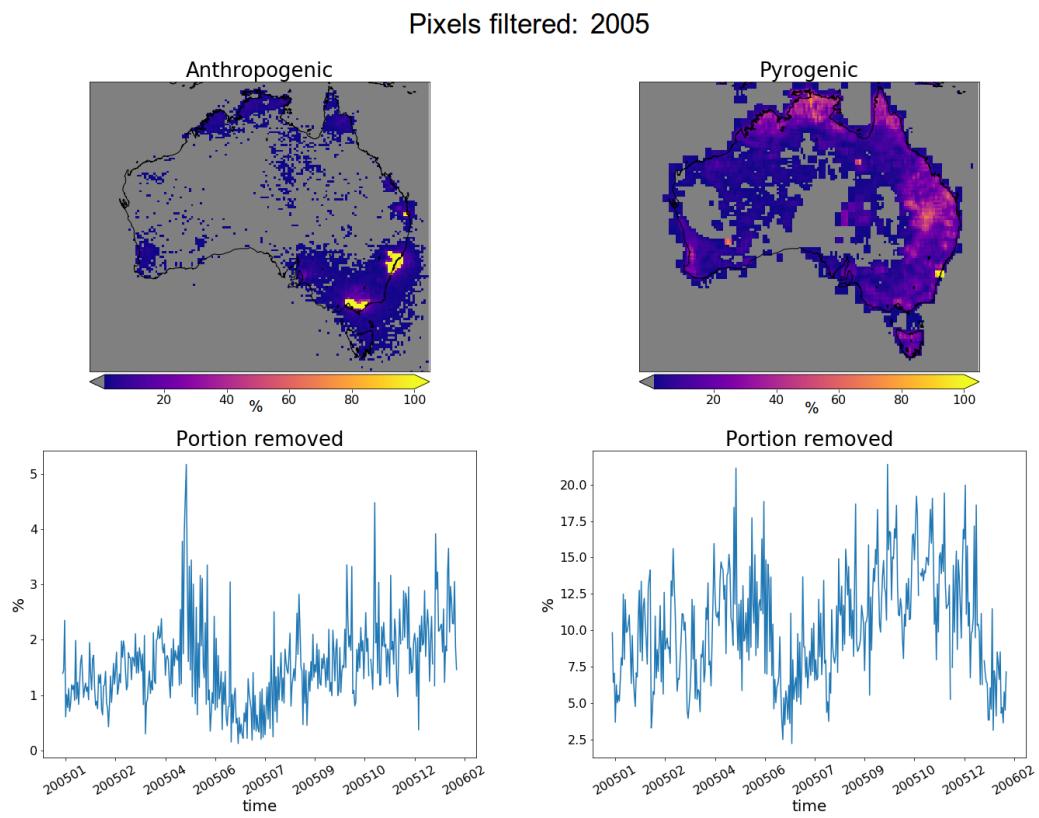


FIGURE 3.2: Top row shows grid squares filtered out by anthropogenic (left) and pyrogenic (right) influence masks over 2005. Bottom row shows portion of Australian grid squares filtered out each day.

4. An AAOD threshold of 0.03 is determined through visual analysis of AAOD distributions over several days, including days with and without influence from active fires, dust, and transported smoke plumes. Grid squares with AAOD over this threshold are considered potentially affected by transported fire smoke.

This method of masking fires can be compared to Marais et al. (2012) and Barkley et al. (2013): Marais et al. (2012) removed pixels colocated with non zero fire counts in any of the prior eight days, within grid squares with  $1^\circ \times 1^\circ$  resolution, and Barkley et al. (2013) used fires from the preceding and current day, within local or adjacent grid squares, at  $0.25^\circ \times 0.3125^\circ$  resolution.

Figure 3.3 shows the effects from filtering HCHO vertical columns with different fire filtering. Vertical column amounts averaged over January, 2005, and the pixel counts are shown side by side. The figure shows the effects of applying an increasingly strict fire filter. Each row has a stricter fire filter applied from top to bottom, with no fire filter on the first row up to filtering pixels from squares with fires up to 8 days prior. Increasing prior days used when creating the fire filter leads to a serious reduction in available pixels, with a wider range of areas dropping to near zero pixel counts. The overall decrease in vertical column HCHO is minor, dropping from  $\sim 4$  to  $\sim 3.9 \times 10^{15}$  molec cm $^{-2}$  between applying no filter and the strictest filter. In this month the fires are most prominent across Queensland and northern NSW, and a large section of the north eastern coast becomes largely filtered out. Increasing the strength of the active fire filter increases the regional impacts on HCHO columns. In this work two prior days of active fires are masked.

Determining the AAOD due to smoke can be difficult since both smoke and dust absorb UV radiation (Ahn, Torres, and Bhartia 2008; Marais et al. 2012). AAOD filtering is designed to remove pixels affected by smoke; however, it may occasionally remove pixels affected by dust. Dust in Australia is highly episodic and false positives in the smoke filter should not affect more than a few days per month, especially over regions with high tree coverage (Shao et al. 2007). The threshold is determined through analysing AAOD over Australia in 4 scenarios: normal conditions, active local fires, transported fire smoke, and large scale dust storms. An example of these scenarios and the AAOD distributions is shown in Figure 3.4. This figure shows AAOD maps and distributions, along with satellite imagery on the same day in column 4 (from <https://worldview.earthdata.nasa.gov/>). The threshold of 0.03 is chosen as it removes very little of a normal day (row 1), and events with higher AAOD show peaks which appear to be distinct from the normal day distribution above this value.

Figure 3.5 shows the extent of pixels filtered out by the pyrogenic filter over 2005 and 2012. One clear hot spot is located over Port Kembla (south of Sydney), most likely due to the flame that burns over the blast furnace stack throughout the year ([https://www.bluescopeillawarra.com.au/community/skylineimages/](http://www.bluescopeillawarra.com.au/community/skylineimages/)). Another hot spot can be seen in Western Australia over Kalgoorlie, where a large open cut gold mine is always open and blasting daily (<https://superpit.com.au/wp-content/uploads/2015/01/Blasting-Information-Sheet.pdf>). In Western Queensland over Mount Isa there is again a mining hotspot, with a blast furnace and several smoke stacks (<http://www.mountisamines.com.au/en/about-mim/Pages/home.aspx>). A large area in southern Queensland/northern NSW is also heavily filtered, potentially due to gas flaring in the Surat Basin, which has thousands of petrol and gas wells (<http://www.ga.gov>.

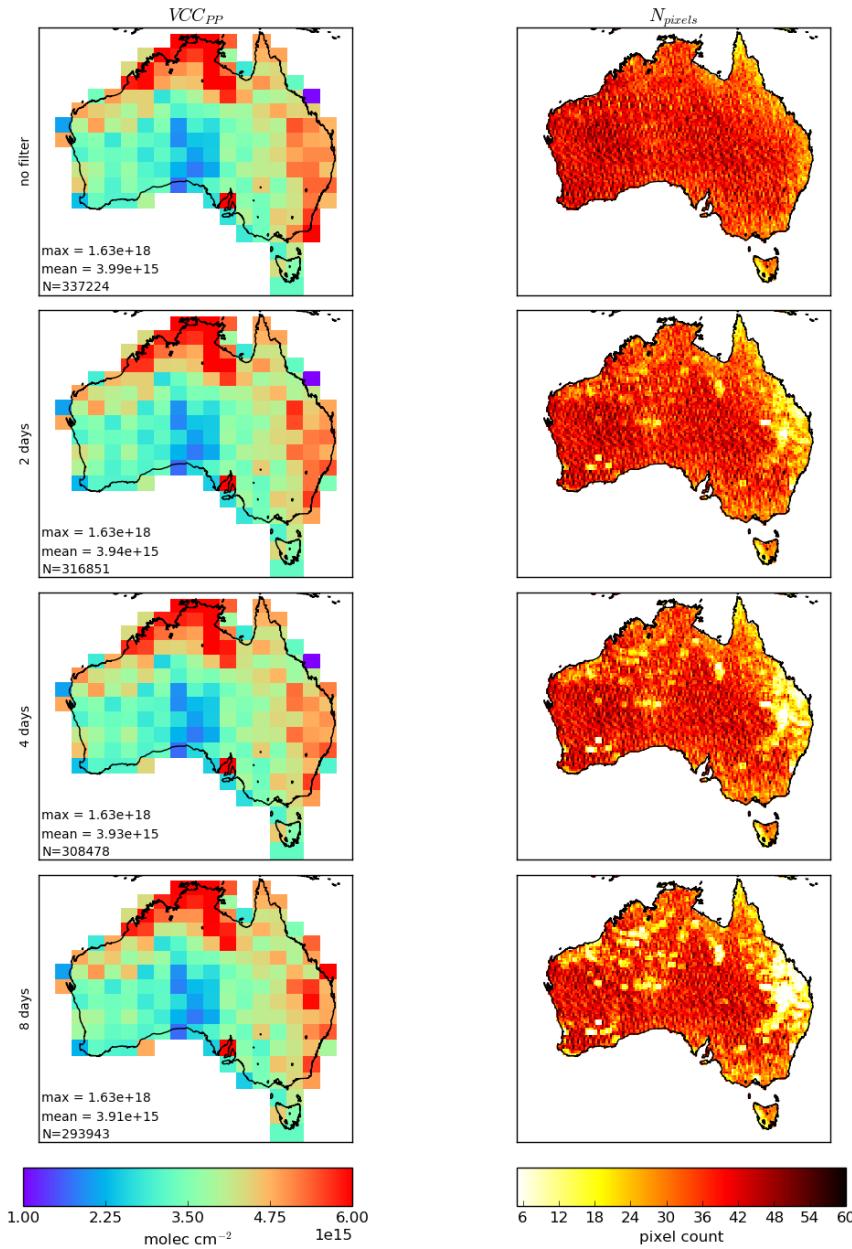


FIGURE 3.3: Recalculated OMI vertical HCHO columns and pixel counts for January 2005. Column 1: VCC<sub>PP</sub> amounts after applying the active fire filter. The filter is applied at  $0.25^\circ \times 0.3125^\circ$  resolution, before output is averaged into  $2^\circ \times 2.5^\circ$  bins. Column 2: Pixel counts after applying the active fire filter (summed into  $0.25^\circ \times 0.3125^\circ$  bins). Row 1-4: increasing number of prior days that have active fires are included when masking fire influence. The first row shows the values without applying any active fire filter. The average and maximum VCC column amount (at  $0.25^\circ \times 0.3125^\circ$  resolution), and number of pixels, is inset as text in column 1 of each row. The second to fourth row respectively show the columns after filtering for adjacent fire on up to 2, 4, and 8 prior days.

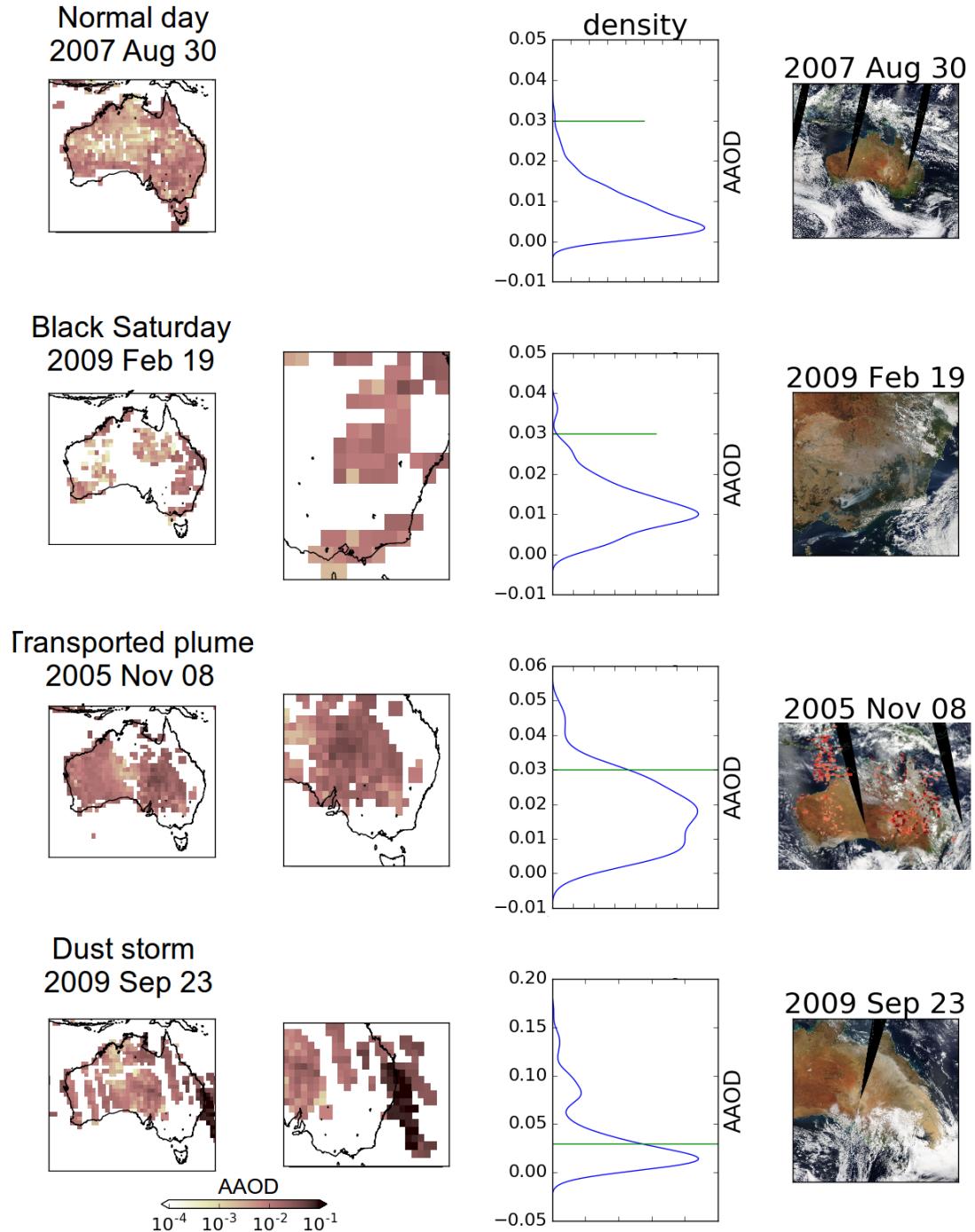


FIGURE 3.4: AAOD from OMAERUVd (columns 1, 2, 3) over Australia for four different scenarios (rows 1-4). Density plots show normalised AAOD frequencies (scale not shown), and the threshold used in this work of 0.03 (see text) is demarcated by a horizontal line. Black Saturday (row 2) refers to the occurrence of widespread bush fires across Victoria. The transported plume in row three can be seen in the overlaid AOD shown in the last column. Scenes in the final column are created using the EOS Worldview website <https://worldview.earthdata.nasa.gov/> from satellite products provided therein.

[au/scientific-topics/energy/resources/petroleum-resources/gas](http://au/scientific-topics/energy/resources/petroleum-resources/gas)). The highest concentrations of removed pixels lie along the northern and eastern coastlines, and correspond with forested areas (see Figure 3.6), which suggests that forest fires are being masked properly. Central Australia is largely unmasked, which could be due to a lack of sufficient vegetation to create fires and smoke visible by satellite. The proliferation of petrol or gas wells (see Figure 3.7) may also lead to AAOD enhancement wherever activity stirs up dust, and could be mistaken as active fires wherever gas flaring occurs. In 2005, 388 gas wells existed in Queensland; however, more than 2000 wells (cumulative) were approved by 2013 (Carlisle 2012). The comparison between fire filtering in 2005 and 2012 (Figure 2.31, bottom plot) does not indicate that the proliferation of gas wells in the intervening years had any strong effect on data filtering.

Additional work was performed to validate the pyrogenic filter through an examination of the relationship between temperature and HCHO (Appendix A). HCHO should be exponentially related to temperature if it is driven by isoprene, with outliers possibly indicative of pyrogenic influence, especially when high emissions are shown on low temperature days. However, fires typically occur on hotter days, and HCHO destruction occurs faster on hotter days, confounding the clear occurrence of outliers (e.g. Zheng et al. 2015). At  $0.25^\circ \times 0.3125^\circ$  resolution, application of the pyrogenic filter slightly strengthens the correlation over the wider south eastern Australian region, with  $r$  increasing from  $\sim 0.31$  to  $\sim 0.38$ . In this thesis the pyrogenic filter is applied at the higher horizontal resolution ( $0.25^\circ \times 0.3125^\circ$ ), and this analysis suggests that the filter should strengthen the relationship between total column HCHO and its biogenic precursors over Australia.

### 3.2.3.2 Anthropogenic filter

Enhanced NO<sub>2</sub> concentrations indicate anthropogenic influence over Australia. We use<sup>mh</sup> NO<sub>2</sub> as a proxy for potential anthropogenic NMVOC emissions, as these could bias the inversion performed in Chapter 3. A filter is designed using the tropospheric NO<sub>2</sub> columns in the OMNO2d product.

NO<sub>2</sub> columns near several major cities in south eastern Australia over 2005 are used to determine a suitable threshold for anthropogenic influence. The mean and standard deviation of tropospheric NO<sub>2</sub> over Australia of measured by satellite is shown in Figure 3.8. Anthropogenic influences are clearly visible near major cities in Australia, and some influence can be seen along nearly every coastline.

The anthropogenic filter is created for each year in two steps:

1. Daily grid squares with NO<sub>2</sub> greater than  $2.0 \times 10^{15}$  molec cm<sup>-2</sup> are flagged as anthropogenic.
2. After taking the yearly average for each grid square, any tropospheric NO<sub>2</sub> columns greater than  $1.5 \times 10^{15}$  molec cm<sup>-2</sup> are flagged for the whole year.

These thresholds are chosen subjectively to ensure the removal of definite anthropogenic influence while not removing too many data points over nearly uninhabited portions of Australia. These thresholds completely remove grid squares over major cities that are likely emitting NMVOCs year round, and also frequently remove grid squares downwind. The effects of applying this filter to the OMNO2d product itself

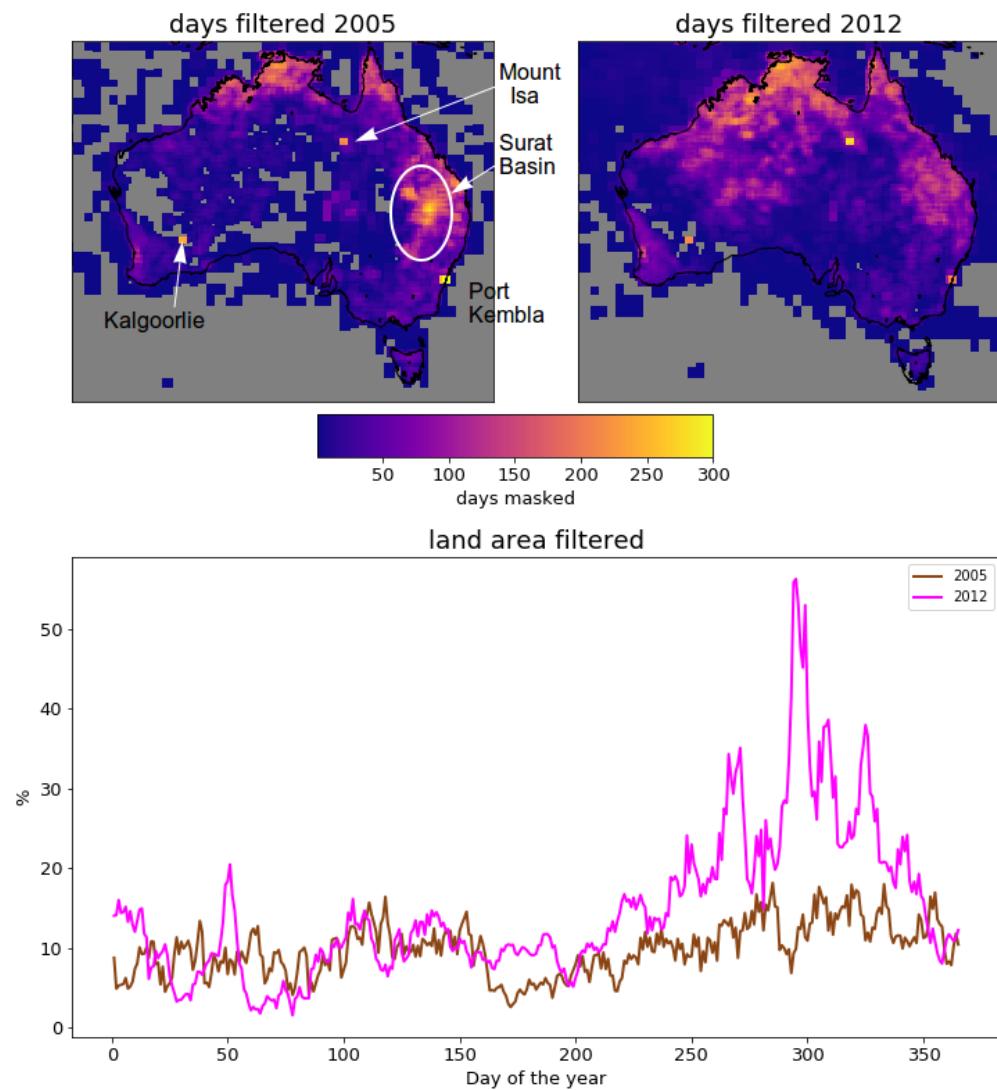


FIGURE 3.5: Top: percentage of pixels filtered out by fire and smoke masks in 2005 (left) and 2012 (right). Bottom: percentage filtered out each day from land squares in Australia for the two years shown.

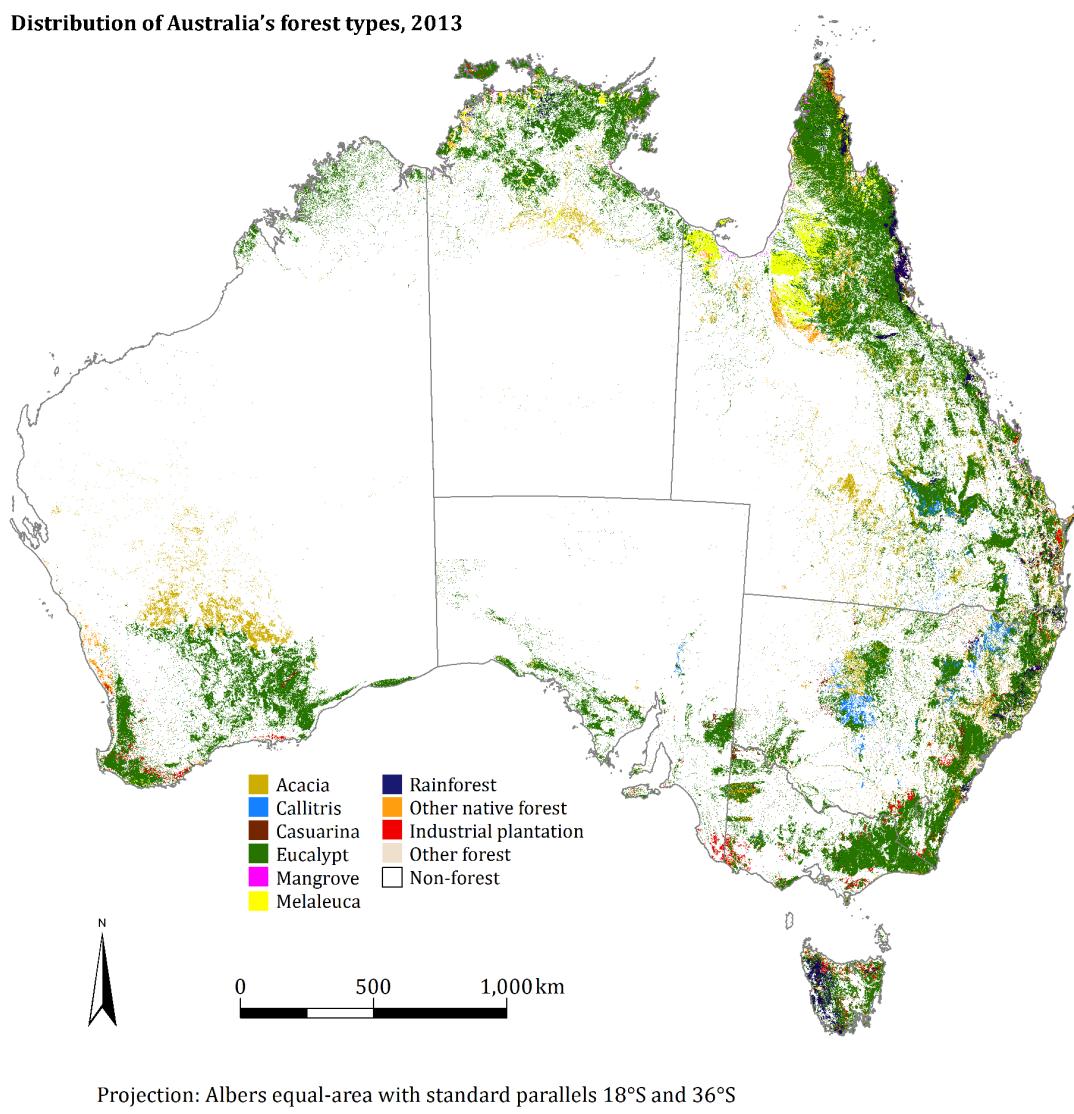


FIGURE 3.6: Forest coverage, coloured by predominant tree species.

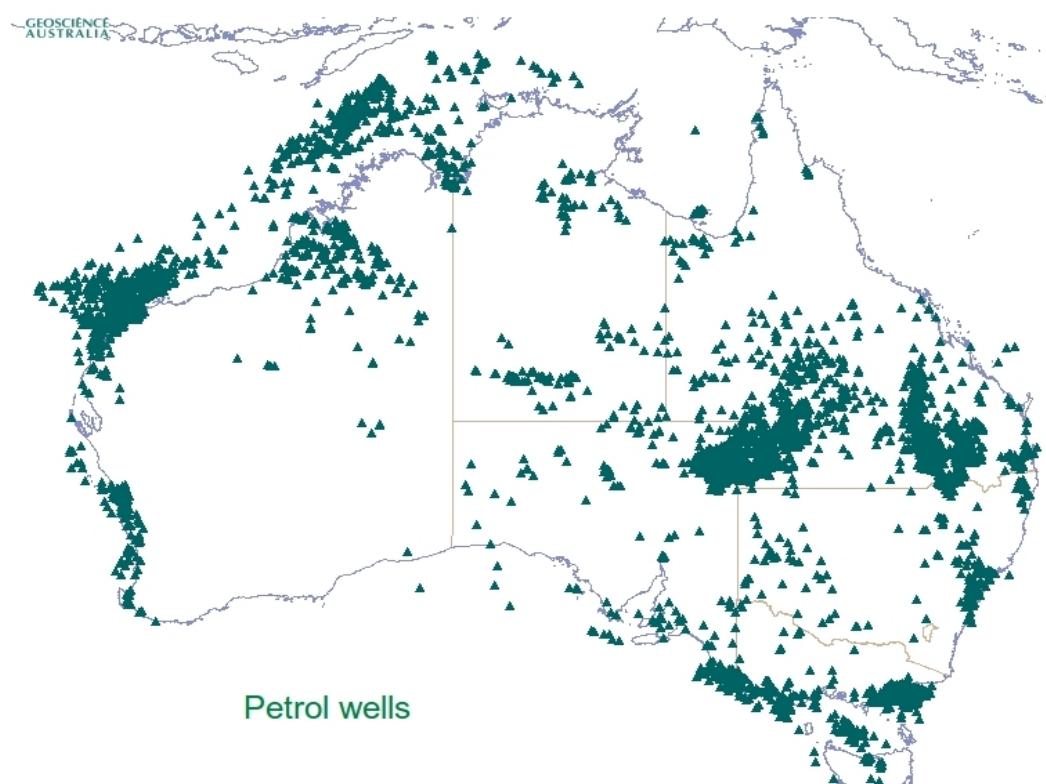
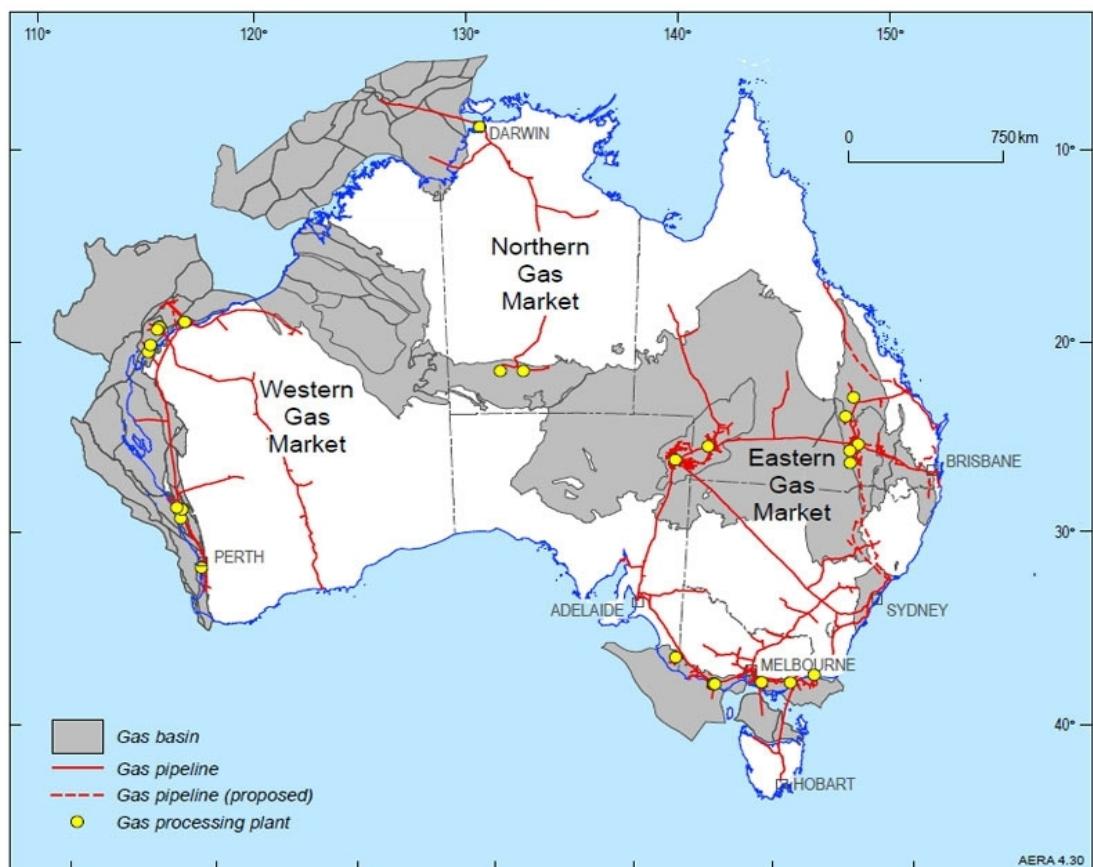


FIGURE 3.7: Top: gas fields and pipelines (2018) for Australia (<http://www.ga.gov.au/scientific-topics/energy/resources/petroleum-resources/gas>). Bottom: petrol Well locations over Australia (as of 2018) (<http://dbforms.ga.gov.au/www/npm.well.search>).

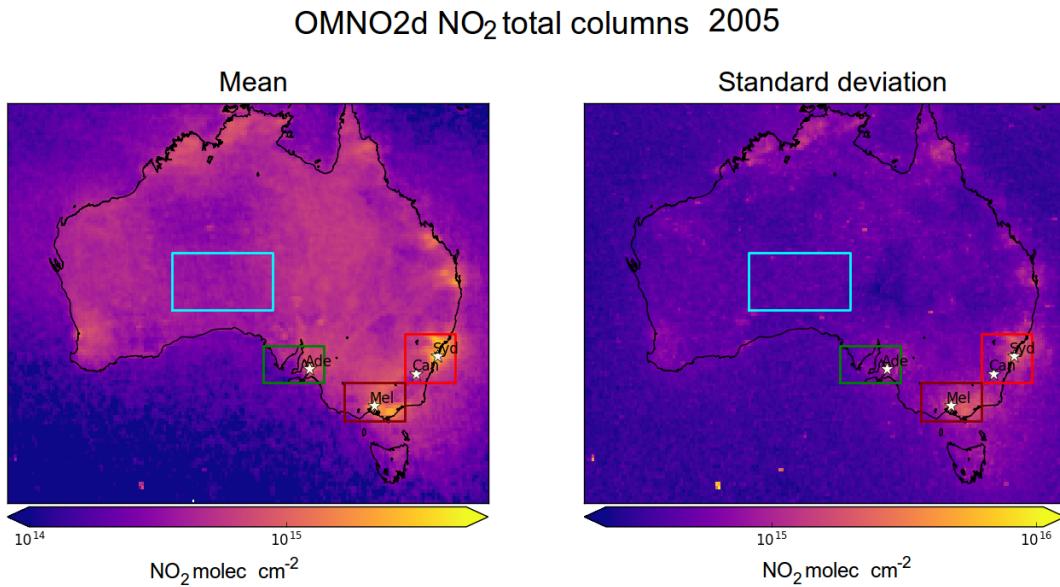


FIGURE 3.8: Mean (left) and standard deviation (right) of OMNO2d daily  $0.25^\circ \times 0.25^\circ$  tropospheric cloud filtered NO<sub>2</sub> columns.

TABLE 3.2: NO<sub>2</sub> averages ( $10^{14}$  molec cm<sup>-2</sup>) by region before and after filtering for anthropogenic emissions using 2005 data from the OMNO2d product.

Region	NO <sub>2</sub>	NO <sub>2</sub> after filtering	% Data lost
Aus	4.9	4.5	2.1%
Sydney	10.4	5.9	25.3%
Adelaide	6.2	5.4	4.0%
Central Aus	3.8	3.7	0.5%
Melbourne	9.6	5.3	25.4%

can be seen in Figure 3.9. Areas over Sydney and Melbourne tend to oscillate around the upper quartile of Australian NO<sub>2</sub> tropospheric columns once the filter is applied. Many coastal regions show some effect from the filtering, and Sydney, Melbourne, and the La Trobe Valley (east of Melbourne) are completely removed.

The same regions as in Figure 3.9 are shown again in Figure 3.10, with NO<sub>2</sub> pixel densities for each region shown, along with the thresholds of  $1.5$  and  $2 \times 10^{15}$  molec cm<sup>-2</sup>. The reduction in NO<sub>2</sub> columns along with the portion of available data removed over Australia and each subset region is listed in Table 3.2. Roughly a quarter of the available data points are removed over Sydney and Melbourne, decreasing the mean NO<sub>2</sub> amounts by  $\sim 50\%$ , while not too much information is lost overall.

### 3.2.3.3 Smearing filter

Smearing is a measure of how much HCHO in a given grid box was produced from isoprene emitted in a different (upwind) grid box. Smearing affects emissions estimates as HCHO enhancements downwind of where precursor emissions occurred

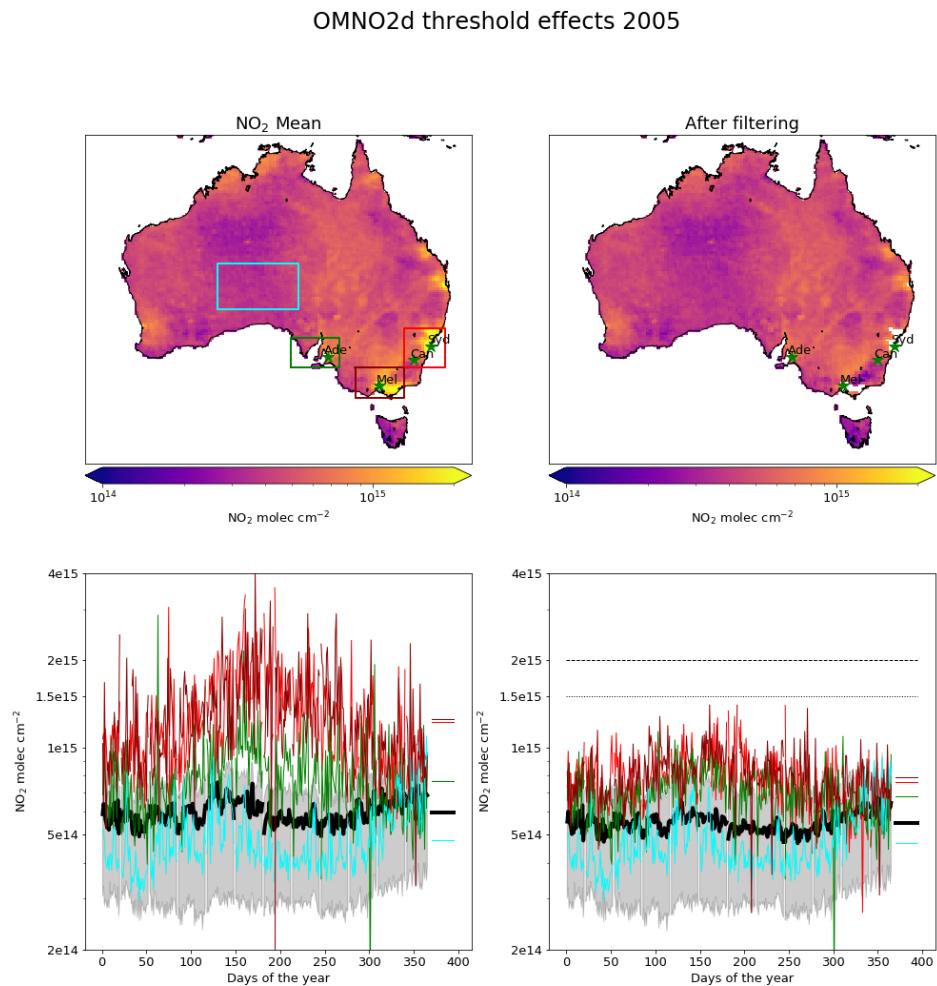


FIGURE 3.9: Top row: 2005 OMNO2d NO<sub>2</sub> column mean before (left) and after (right) applying the anthropogenic threshold filters as described in the text. Bottom row: time series for Australia, and each region (by colour) shown in the bottom panels, with mean for that region shown as as horizontal bars within the bottom panels at the right (over 360-400 on the x-axis). The time series before and after applying the anthropogenic filter are shown on the left and right respectively. The thicker black line shows the mean value over all of Australia, and the grey shaded area shows the inter-quartile range. The dashed and dotted horizontal lines show daily and yearly mean thresholds respectively.

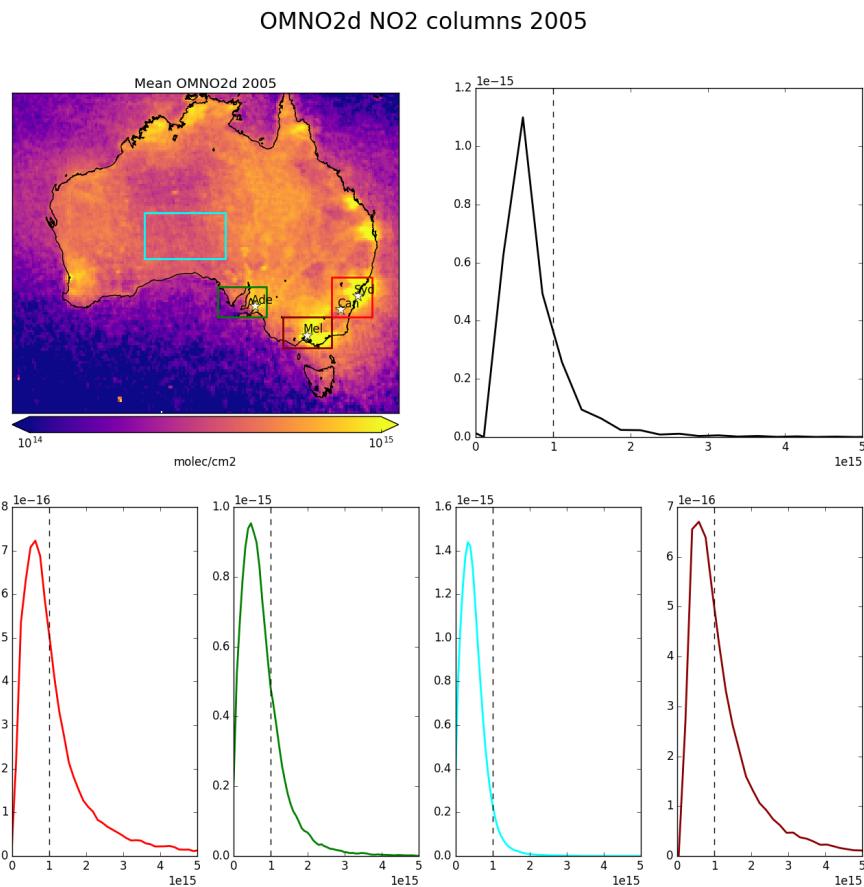


FIGURE 3.10: 2005 OMNO2d NO<sub>2</sub> column means (top left) and distributions (top right) for Australia, and each region shown in the area map (by colour). Vertical dashed lines show the threshold for anthropogenic influence; any columns above this value are filtered out. The vertical axis is normalised so that area under the curve adds to unity, and as such is not important except as a visual measure of the relative width between the distributions.

lead to misinterpretation of local emissions. In high  $\text{NO}_x$  ( $>\sim 1 \text{ ppb}$ ) environments, isoprene has a lifetime on the order of 30 minutes, and HCHO can be used to map isoprene emissions with spatial resolution from 10-100 km (Palmer et al. 2003). In low  $\text{NO}_x$  conditions, isoprene has a longer lifetime (hours) and may form HCHO further from the source area (Fan and Zhang 2004; Liu et al. 2016b; Liu et al. 2017b). Over Australia,  $\text{NO}_x$  levels are generally low and smearing is therefore expected to be important. Smearing limits the horizontal resolution of the linear top-down inversion process, as a finer resolution increases sensitivity to transport. Horizontal transport *smears* the HCHO signal so that its source location would need to be calculated using wind speeds and loss rates (Palmer et al. 2001; Palmer et al. 2003). In this chapter smearing affected grid squares are filtered out prior to application of Equation 3.10.

**3.2.3.3.1 Calculation of smearing** Smearing has been analysed in several publications (e.g., Martin et al. 2003; Palmer et al. 2003; Millet et al. 2006; Stavrakou et al. 2009; Marais et al. 2012; Barkley et al. 2013; Zhu et al. 2014b; Wolfe et al. 2016; Surl, Palmer, and Abad 2018) and is often calculated using the method used in this thesis, as first described by Palmer et al. (2003). This involves using two model runs, one of which has isoprene emissions scaled globally by a constant (generally from 0.5 to 2). From Section 3.2.5, Equation 3.10 states that the modelled slope ( $S$ ) is the yield of HCHO per C of emitted isoprene divided by the HCHO loss rate ( $S = \frac{Y_{\text{isop}}}{k_{\text{HCHO}}}$ ). Using two runs of GEOS-Chem with differing isoprene emissions but otherwise identical *provides we have<sup>mh</sup>*:

$$\begin{aligned} \text{Run}_1 : \Omega_{\text{HCHO}} &= SE_{\text{isop}} + \Omega_0 \\ \text{Run}_2 : \Omega'_{\text{HCHO}} &= S'E'_{\text{isop}} + \Omega'_0 \end{aligned} \quad (3.2)$$

There are several assumptions that need to be understood, as these are what is tested by the smearing calculation. The initial assumption is that the system is at steady state, with no transport of isoprene affecting HCHO columns. This is the basis for equation 3.2. It is also assumed that background values ( $\Omega_0$ ) are from oxidation of methane and other long-lived VOCs, so that  $\Omega_0 = \Omega'_0$ . Between these two runs *only the E term is changed-we are only changing the E term<sup>mh</sup>*. Chemistry is unchanged so that the yield and loss rate should not change between the two runs:

$$S = S' = \frac{Y_{\text{isop}}}{k_{\text{HCHO}}} \quad (3.3)$$

Equations 3.2 may then be combined as follows:

$$\begin{aligned} \text{Run}_1 - \text{Run}_2 : \Omega_{\text{HCHO}} - \Omega'_{\text{HCHO}} &= SE_{\text{isop}} - S'E'_{\text{isop}} + \Omega_0 - \Omega'_0 \\ \Omega_{\text{HCHO}} - \Omega'_{\text{HCHO}} &= S(E_{\text{isop}} - E'_{\text{isop}}) \\ \Delta\Omega_{\text{HCHO}} &= S\Delta E_{\text{isop}} \\ \hat{S} &\equiv \frac{\Delta\Omega_{\text{HCHO}}}{\Delta E_{\text{isop}}} \approx \frac{Y_{\text{isop}}}{k_{\text{HCHO}}} \end{aligned} \quad (3.4)$$

TABLE 3.3: Smearing filters or typical slopes ( $S$ ) from literature.

Source	min. (s)	max. (s)	type <sup>a</sup>	Region
Palmer et al. (2003)	1270	2090	Range	North America <sup>b</sup>
Marais et al. (2012)		4000	Limits	Africa
Barkley et al. (2013) <sup>c</sup>	1300	1800	Limits	South America
Surl, Palmer, and Abad (2018)	2200	4900	Range	India
This Thesis	800	4600	Limits	Australia

a: Slope *ranges* are observed or modelled  $S$ , while smearing *limits* are the applied acceptable limits for  $S$ .

b: Slope range for summer only.

c: Assumed HCHO lifetime of 2.5 hours implies yields from 0.14 to 0.2 per C, consistent with box modelling.

This allows the combination of outputs from the two runs to determine where  $\hat{S}$  diverges from expected values for  $S$ .

Potential smearing is masked by checking a daily modelled value for  $\hat{S} \approx Y_{isop} / k_{HCHO}$  against thresholds. By assuming that midday HCHO lifetime ( $\tau = 1/k_{HCHO}$ ) typically falls within 1.5 to 4 hrs (as seen in the USA; Palmer et al. (e.g., 2006) and Wolfe et al. (2016)) and isoprene-to-HCHO yield (HCHO per isoprene carbon emitted) lies within the range 0.2 to 0.4 (scenarios estimated in Palmer et al. (2003)), one can set a simple bound on  $\hat{S}$  of  $[0.2 \times 1.5, 0.4 \times 4]$  hrs or 1080 to 5760 seconds. As  $\text{NO}_x$  levels across Australia are relatively low, and lower  $\text{NO}_x$  levels reduce the prompt yield (Palmer et al. 2003; Wolfe et al. 2016). In this work the threshold range is the simple bound shown here, reduced by 20% and rounded to the nearest hundred leading to a bounding range of 800 to 4600 for  $\hat{S}$ . This range strikes a balance between unlikely modelled yields and the amount of data lost to filtering. Table 3.3 compares the smearing filter for Australia used in this thesis to typical values used in previous work for other regions.

GEOS-Chem is run with normal isoprene emissions and with isoprene emissions halved, then Equation 3.4 ( $\hat{S} = \frac{\Delta \Omega_{HCHO}}{\Delta E_{isop}}$ ) provides  $\hat{S}$ . Here  $\Delta$  represents the difference (daily 1300-1400 local time) between default and scaled runs. If  $\hat{S}$  sits outside the 800-4600 range then ~~we remove<sup>mh</sup>~~ that grid square day is removed<sup>mh</sup> from both  $S$  and subsequent a posteriori emission calculations. A relatively large change in  $\Omega_{HCHO}$  compared to local emissions ( $\hat{S} > 4600$ ) suggests HCHO production is from non-local isoprene emissions. Alternatively, a relatively low value of  $\hat{S}$  ( $\hat{S} < 800$ ) suggests emissions from the local grid square are being exported before they form HCHO.

**3.2.3.3.2  $\text{NO}_x$  dependence**  $\text{NO}_x$  concentration directly affects the fate of VOCs in the atmosphere, influencing HCHO production by isoprene and therefore the amount of smearing. In low  $\text{NO}_x$  environments, reported HCHO yields from isoprene are around 0.2 - 0.3 C per C (or 100-150 molar %), while in high  $\text{NO}_x$  environments this value becomes two to three times higher (Palmer et al. 2003; Wolfe et al. 2016). Some values for HCHO yield from prior literature are shown in Table 3.4.

TABLE 3.4: Isoprene to HCHO yields and lifetime.

HCHO Yield (molar % )	Lifetime	NO <sub>x</sub> background	Source
315±50		High	a
285±30		High	a
225	35 min	High	b
450		High	c
235		1 ppbv	d
150		Low	b
150		Low	c
150		0.1 ppbv	d

a Atkinson and Arey (2003): Table 2, Yield from Isoprene reaction with OH, two values are from two referenced papers therein.

b Palmer et al. (2003): lifetimes assume [OH] is 1e15 mol cm<sup>-3</sup>.

c Wolfe et al. (2016): “prompt yield”: change in HCHO per change in isop<sub>0</sub>.  $[isop]_0 = [isop] \exp(k_1[OH]t)$ ; where  $k_1$  is first order loss rate. Effectively relates HCHO abundance with isoprene emission strength.

d Dufour et al. (2008): One-day yields from oxidation modelled by CHIMERE, using MCM reference scheme.

The effect of NO<sub>2</sub> on smearing can be seen in Figure 3.11. This plot shows how smearing over Australia compares to satellite NO<sub>2</sub>, with smearing distributions binned by NO<sub>2</sub> both with and without filtering for smearing. At lower NO<sub>2</sub> levels smearing is more frequently below the lower threshold. These low values decrease in frequency, and have less affect on the mean  $\hat{S}$  at around  $5 \times 10^{14}$  molec cm<sup>-2</sup> NO<sub>2</sub>.

### 3.2.4 GEOS-Chem emissions

In this work MEGAN (version 2.1) is run as a module within GEOS-Chem (version 10.01) (see Section 2.3.4). The chemical model is driven by GEOS-5 meteorological fields at 0.5°x 0.666° horizontal resolution. GEOS-Chem output is averaged onto 47 vertical levels at 2° × 2.5°, based on chemistry and transport calculated every 30 and 15 minutes respectively. Isoprene emissions from the default *tropchem* simulation are referred to as the a priori emissions. When shown as part of a formula the a priori emissions are denoted as  $E_{GC}$ .

### 3.2.5 Relationship between isoprene emissions and formaldehyde

Tropospheric HCHO production is primarily due to the oxidation of VOC precursor species ( $VOC_i$ ). Background concentrations are driven by methane; a longer lived ( $\sim 8$  yr) VOC. Over continental land masses, the variability in HCHO is driven by shorter lived precursor emissions (Chance et al. 2000; Palmer et al. 2003). HCHO is

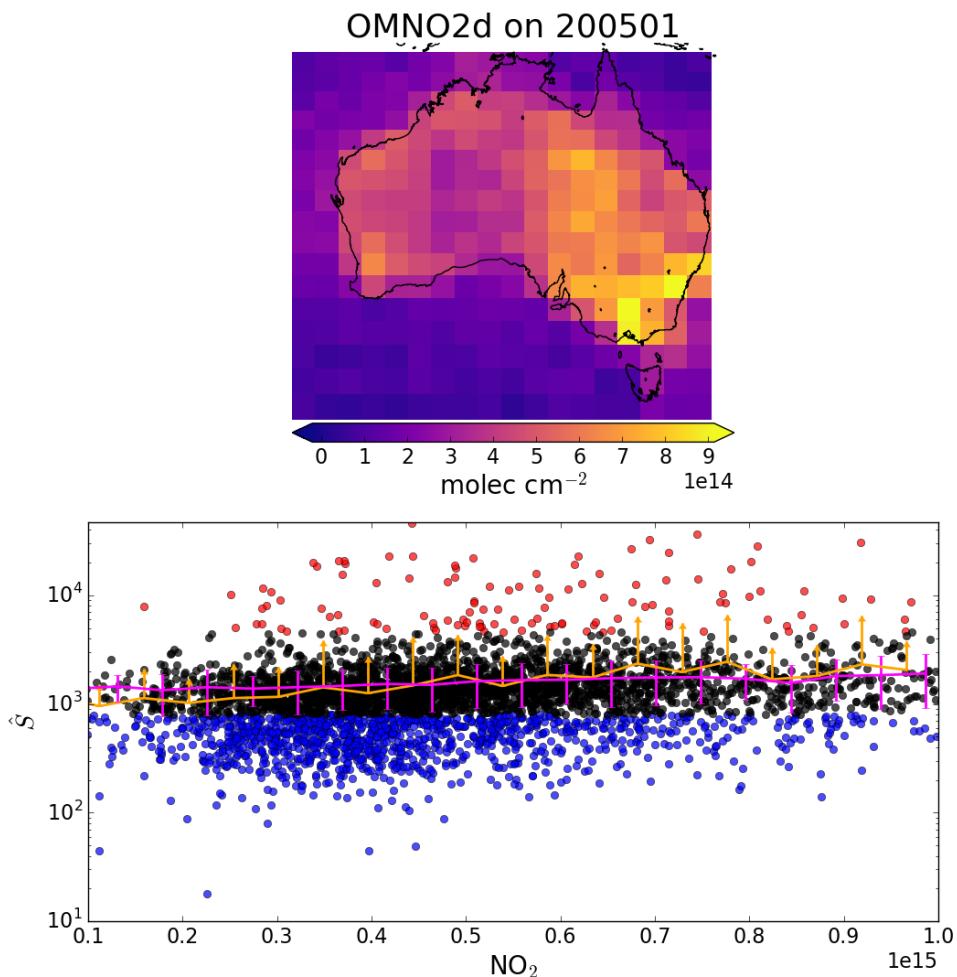
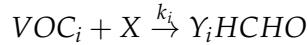


FIGURE 3.11: Top: OMNO2d tropospheric NO<sub>2</sub> columns averaged into  $2^\circ \times 2.5^\circ$  horizontal bins for Jan, 2005. Bottom: Scatter plot of NO<sub>2</sub> against smearing calculations from GEOS-Chem ( $\hat{S}$ ), with points above and below the smearing threshold range of 800-4600 s coloured red and blue respectively. Points are binned by NO<sub>2</sub> with and without the smearing filter applied (orange and magenta respectively). Overplotted is the mean and standard deviation (error bars) within each bin. Due to the logarithmic Y scale **only the positive direction of standard deviations is shown—we only show the positive direction of standard deviations<sup>mh</sup>** for unfiltered data.

then produced quickly from short-lived intermediates:



where  $X$  is an oxidant,  $Y_i$  is HCHO yield (per C atom in  $VOC_i$ ), and  $k_i$  is the reaction rate constant. In specific conditions described below, HCHO total columns ( $\Omega$ ; molec  $\text{cm}^{-2}$ ) can be linearly related to isoprene emissions. In Australia the effective molar HCHO yield from isoprene has not been extensively studied, while in other continents this value varies from 1-3 depending on local  $\text{NO}_x$  concentrations (e.g., Palmer et al. 2006; Millet et al. 2006; Bauwens et al. 2016; Surl, Palmer, and Abad 2018).

The isoprene to HCHO relationship is derived using several assumptions that are outlined here. The first assumption is that HCHO is at steady state, which implies production ( $P_{HCHO}$ ) and loss ( $L_{HCHO}$ ) are equivalent:

$$\frac{d\Omega}{dt} = 0 = P_{HCHO} - L_{HCHO} \quad (3.5)$$

This is reasonable during midday when isoprene emissions are steady and  $\Omega$  has had time to stabilise. The second assumption is that loss (with loss rate constant  $k_{HCHO}$ ) is only first order, such as from photolysis and oxidation:

$$L_{HCHO} = k_{HCHO}\Omega \quad (3.6)$$

This assumption means that loss due to transport must be negligible as it is not first order. This assumption is reasonable for large enough grid boxes as transport becomes negligible relative to the linear (first order) terms. Production and loss are on the order of minutes, and grid box sizes in this work are rectangular with  $\sim 200 \text{ km}$  edge lengths. Monthly averaged wind speeds rarely exceed  $20 \text{ km hr}^{-1}$  over Australia, meaning HCHO and precursor transport remain minor. Transport can still be an issue, however, and is handled by applying a smearing filter described in Section 3.2.3.3.

Another assumption is that  $\Omega$  production above the background level is due only to precursor emissions ( $E_i$ ; atoms  $\text{C cm}^{-2}\text{s}^{-1}$ ) multiplied by their yields to HCHO ( $Y_i$ ):

$$P_{HCHO} = \sum_i Y_i E_i \quad (3.7)$$

By combining Equations 3.5, 3.6, and 3.7,  $\Omega$  can be related to [we can relate  \$\Omega\$  to  \$\text{m}^h\$](#)  precursor emissions:

$$\begin{aligned} k_{HCHO}\Omega &= \sum_i Y_i E_i \\ \Omega &= \frac{1}{k_{HCHO}} \sum_i Y_i E_i \end{aligned} \quad (3.8)$$

Finally, it is assumed isoprene emissions are driving changes in  $\Omega$  (as assumed elsewhere, e.g., Palmer et al. 2003; Millet et al. 2008; Marais et al. 2014; Stavrakou et al.

2015) and lump other terms together:

$$\sum_i Y_i E_i = Y_{isop} E_{isop} + \sum_{i \neq isop} Y_i VOC_i \quad (3.9)$$

This assumption is reasonable only over continental land masses, and can be false when pyrogenic or anthropogenic emissions influence the HCHO column. These scenarios are filtered to remove pyrogenic and anthropogenic influence using independent satellite measurements (see Section 3.2.3). The linear relationship between isoprene emissions and  $\Omega$  is determined by equating  $P_{HCHO}$  and  $L_{HCHO}$  from Equations 3.7 and 3.6, substituting Equation 3.9, and assuming that the lumped terms make up the background:

$$\begin{aligned} k_{HCHO}\Omega &= Y_{isop} E_{isop} + \sum_{i \neq isop} Y_i VOC_i \\ \Omega &= \frac{Y_{isop}}{k_{HCHO}} E_{isop} + \Omega_0 \\ &= S \times E_{isop} + \Omega_0 \end{aligned} \quad (3.10)$$

Here  $S$  is the slope of the regression between  $E_{isop}$  and  $\Omega$ :  $S \equiv \frac{Y_{isop}}{k_{HCHO}}$ .

### 3.2.6 Calculation of modelled slope

To determine  $S$ , the link between biogenic isoprene and midday column HCHO, GEOS-Chem is used. The term  $E_{GC}$  is used when discussing isoprene emissions estimated within GEOS-Chem and  $\Omega_{GC}$  is used to represent simulated HCHO total column. The method to calculate  $S$  using GEOS-Chem follows roughly the following three stages:

1. Hourly gridded model outputs  $E_{GC}$  (atoms C cm<sup>-2</sup> s<sup>-1</sup>), along with  $\Omega_{GC}$  (molec cm<sup>-2</sup>) at 13:30 LT daily are extracted from the biogenic-emissions-only run.
2. Filtering removes gridded output on days where grid squares are likely to be affected by smearing (see Section 3.2.6.1).
3. A reduced major axis regression slope is determined between  $\Omega_{GC}$  and  $E_{GC}$  for every month of modelled output (one value per day) for each grid square (Figure 3.12).

Modelled background concentrations can be ignored here as they do not affect slope calculation. This effectively provides the monthly gridded slope ( $S$ ) between biogenic isoprene emissions and HCHO columns, in units of seconds.

Figure 3.12 (top left) shows how  $S$  varies spatially over Australia for an example mid-summer month. High  $S$  values suggest high sensitivity of HCHO to isoprene emissions. Some areas can be seen to be very sensitive to emissions, such as the west coast and Eyre basin, which is likely due to the low isoprene and HCHO levels in those areas. The regression coefficients also vary spatially (top right), and some areas show little correlation. It is likely that this is due to weather, transport, and a lack of local emission sources. The slopes in the bottom panel show a small sample of scatter and regression plots. These can range widely due to differences in emission

and yield parameters, which plays a role in the smearing filters described in Section 3.2.3.3. Due to the  $2^\circ \times 2.5^\circ$  horizontal resolution of GEOS-Chem, calculations over coastal grid boxes that are mostly oceanic are often discarded as the change in HCHO is not dominated by emissions of isoprene, as is assumed for equation 3.10.

### 3.2.6.1 Smearing

One issue with slope calculation is potential transport (also known as smearing), either of isoprene transported in from outside the local grid box (before any HCHO is formed), or of HCHO formed by local emissions but transported out of the local grid box. A *smearing* filter is created (see Section 3.2.3.3) which filters out grid squares on days when transport is likely. Days where smearing is expected to affect local levels of HCHO are removed before calculating  $S$ , and a simple analysis is performed on how the filter affects monthly slope, correlation, and uncertainty. Figure 3.13 shows the calculated slope for 2005-2012, along with its 95% confidence interval for the  $2^\circ \times 2.5^\circ$  grid box containing Sydney. The monthly and multi-year monthly averages are shown before and after the smearing filter is applied. The filter slightly reduces the amplitude of the seasonal cycle, raising the January minimum and lowering the June and July maximum. Filtering slightly improves the correlation coefficient throughout the year. This is likely due to smearing being prevalent in areas where the relationship between a priori emissions and HCHO columns is already be weak (due to low emissions or unsuitable meteorological conditions). More data are filtered in summer, presumably due to higher biogenic isoprene emissions over summer, making transport more noticeable on windy days. Anthropogenic precursor emissions are not halved between the two runs, and they make up relatively more of the total HCHO precursor concentration in winter (when biogenics are lower). This may lower the smearing filter signal over Sydney and other densely populated areas (especially in winter). This plot has been repeated for several grid squares over Australia (not shown) sometimes showing more filtering throughout winter months, and sometimes over summer months. When calculating top-down emissions the smearing-filtered slope ( $S$ ) is generally used for each grid square month. The multiple year monthly averaged slope is used instead when the regression coefficient ( $r$ ) is less than 0.4, or the number of data points used in the regression ( $n$ ) is less than 10. When  $r$  for the multiple year slope is also lower than 0.4 (does not happen in the example grid square), no estimation is performed. This happens only infrequently, and only in locations with very limited isoprene emissions such as the lake Eyre basin.

Smearing can be dependent on local or regional weather patterns, as greater wind speeds will reduce the time any emitted compound stays within the local grid box. As such smearing can vary both spatially and temporally. Smearing is also sensitive to time of day, season, and latitude, as lower insolation results in slower photolysis. Figure 3.14 shows smearing and how frequently grid squares are filtered using the smearing filter. The smearing filter is more active in winter and spring, especially at higher latitudes. Grid squares along the east coast are filtered frequently in all seasons, as well as grid squares in the high smearing lake Eyre region in northeastern South Australia. Data loss from smearing is approximately 30% over the entirety of Australia during summer, and 40% during winter. The data loss is generally higher towards the north east and southern coastlines, and at lake Eyre.

HCHO trop column vs isoprene emissions Jan, 2005

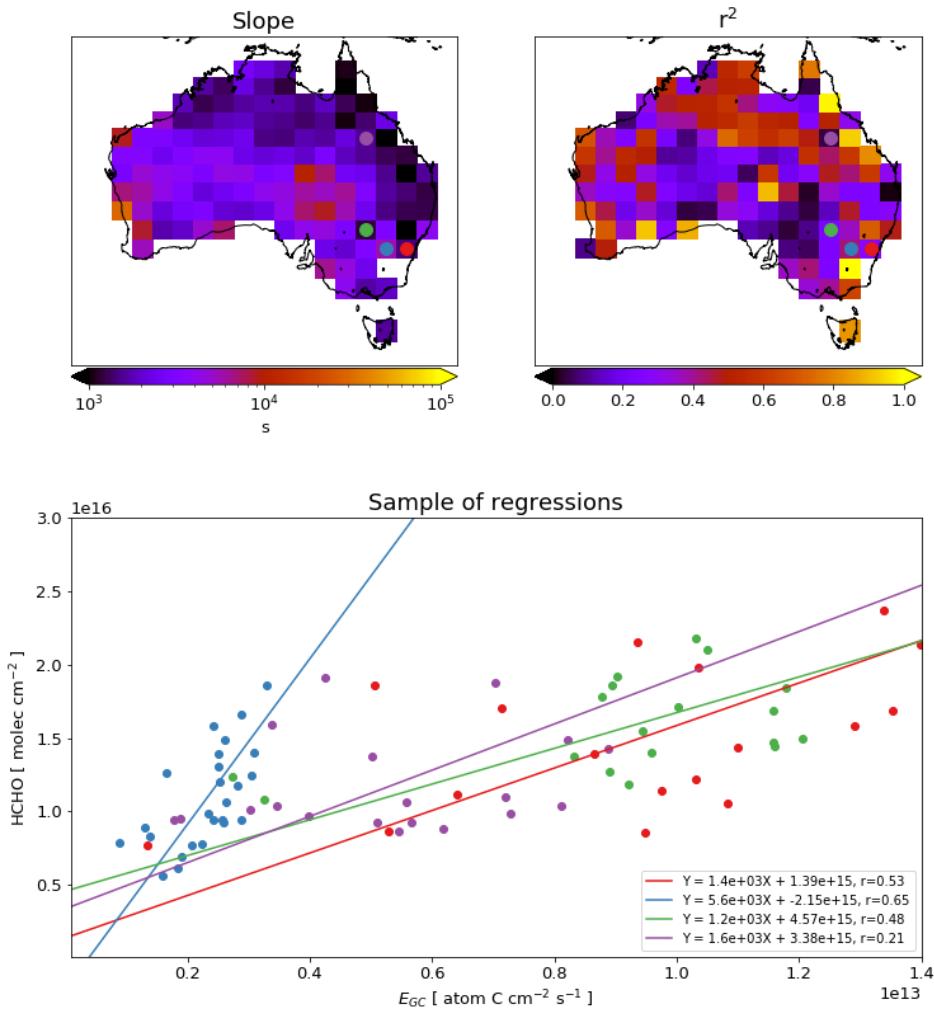


FIGURE 3.12: Top left: reduced major axis slope between modelled tropospheric column HCHO and isoprene emissions ( $E_{GC}$ ) using mid-day (13:00-14:00 LT) values over for January 2005, per grid square at  $2^\circ \times 2.5^\circ$  horizontal resolution. Top right: Squared reduced major axis correlation coefficient for regression in top left. Bottom: Sample of correlations from four grid squares. Coloured dots in top panels correspond to the colour of the regressions shown in bottom panel.

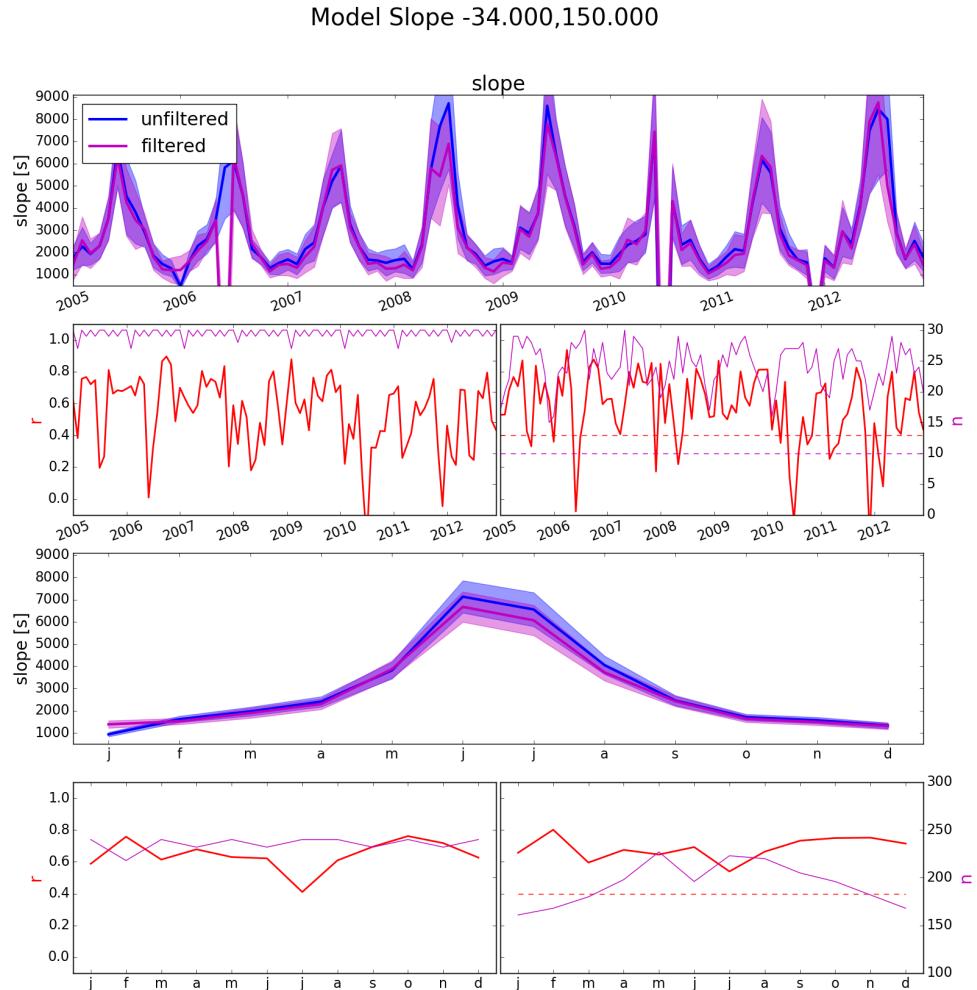


FIGURE 3.13: Row 1: monthly slope along with 95% confidence interval both before (blue) and after (magenta) applying the smearing filter for the model grid square containing Sydney over 2005-2012. Row 2: regression coefficient and data point counts before (left) and after (right) performing filtering for slopes shown in row 1. Additionally, limits for  $r$  and  $n$  used in slope utilisation (see text) are shown with dashed lines. Row 3: slope and confidence interval using the multi-year dataset for each month. Row 4: regression coefficient and data point counts for row 3.

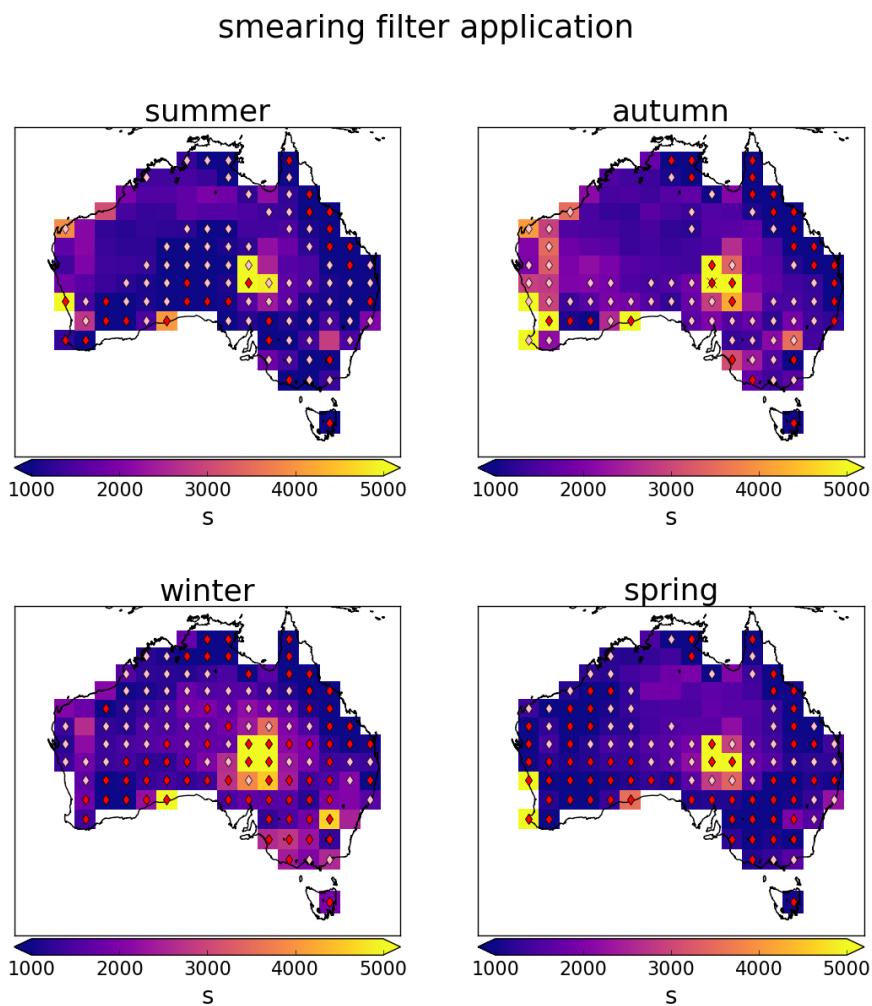


FIGURE 3.14: Seasonally averaged smearing ( $\hat{S}$ , see text) over 2005. Diamonds represent grid squares which have had at least 10 (pink) or 30 (red) days removed due to the smearing filter over the season. Red crosses show where the filter has removed all data.

### 3.2.7 Modelled background HCHO

There are two simple ways to determine the modelled background HCHO, one of which involves running the model with no isoprene emissions. Since variation in HCHO columns is assumed to only depend on isoprene emissions, `theourmh` background term is theoretically identical to the simulated HCHO without isoprene emissions. The other method uses HCHO over the remote Pacific Ocean at matching latitudes and times, which emulates how the background is determined for the satellite measured HCHO. Figure 3.15 shows the background total column HCHO calculated in these two different manners, and how they compare to each other and normal levels for an example month (January, 2005). The difference between these definitions is approximately 15%, and is mostly caused by non-isoprene HCHO precursors (such as monoterpenes). Non-isoprene HCHO precursor emissions are neglected in the top-down estimation technique used in this work, which introduce some uncertainty (Section 3.4.2). Generally in summer months the continental HCHO levels are over 3 times higher than background levels, so the choice of background definition has only a very small impact on the final results. For consistency with the satellite data, backgrounds are determined using the remote Pacific. Background HCHO for any latitude in this thesis is calculated by averaging longitudinally (140°W to 160°W) the matching latitudes over the remote Pacific.

### 3.2.8 Calculation of Emissions

Top-down emissions estimates are calculated using OMHCHO (see Section 2.4) slant columns and an updated AMF calculated using code written by Paul Palmer and Randal Martin, with modifications by Luke Surl (see Section 2.6.3.2). In Chapter 2 there are three calculations of the AMF referred to as  $AMF_{OMI}$  (the original OMI AMF),  $AMF_{GC}$  (AMF recalculated with GEOS-Chem shape factors), and  $AMF_{PP}$  (AMF recalculated using GEOS-Chem shape factors and scattering weights). The new emissions are calculated (using the  $AMF_{PP}$ ) are referred to as the a posteriori from here onward, or  $E_{OMI}$  in formulae.

A posteriori emissions are calculated using the linear relationship described in Section 3.2.5 using the modelled slope  $S$  calculated in the prior section and satellite HCHO columns recalculated in 2.6:

$$\begin{aligned}\Omega_{OMI} &= S \times E_{OMI} + \Omega_0 \\ E_{OMI} &= \frac{\Omega_{OMI} - \Omega_0}{S}\end{aligned}\tag{3.11}$$

This is the same as equation 3.10, except now the satellite HCHO ( $\Omega_{OMI}$  and  $\Omega_0$ ) is used.  $\Omega_0$  is calculated using  $\Omega_{OMI}$  in the remote Pacific averaged monthly and longitudinally, for each latitude. This leaves  $E_{OMI}$  as the only unknown once the satellite measurements are processed to match the temporal and horizontal resolution of  $S$ .

One potential issue in this top-down estimation technique is the low number of valid satellite measurements that may occur due to the higher solar zenith angles in winter and at higher latitudes. Another issue is that negative emissions are commonly calculated in areas wherever measured HCHO columns are lower than background amounts (as  $E_{OMI} = \frac{\Omega_{OMI} - \Omega_0}{S}$ ) due to uncertainty in the background calculation.

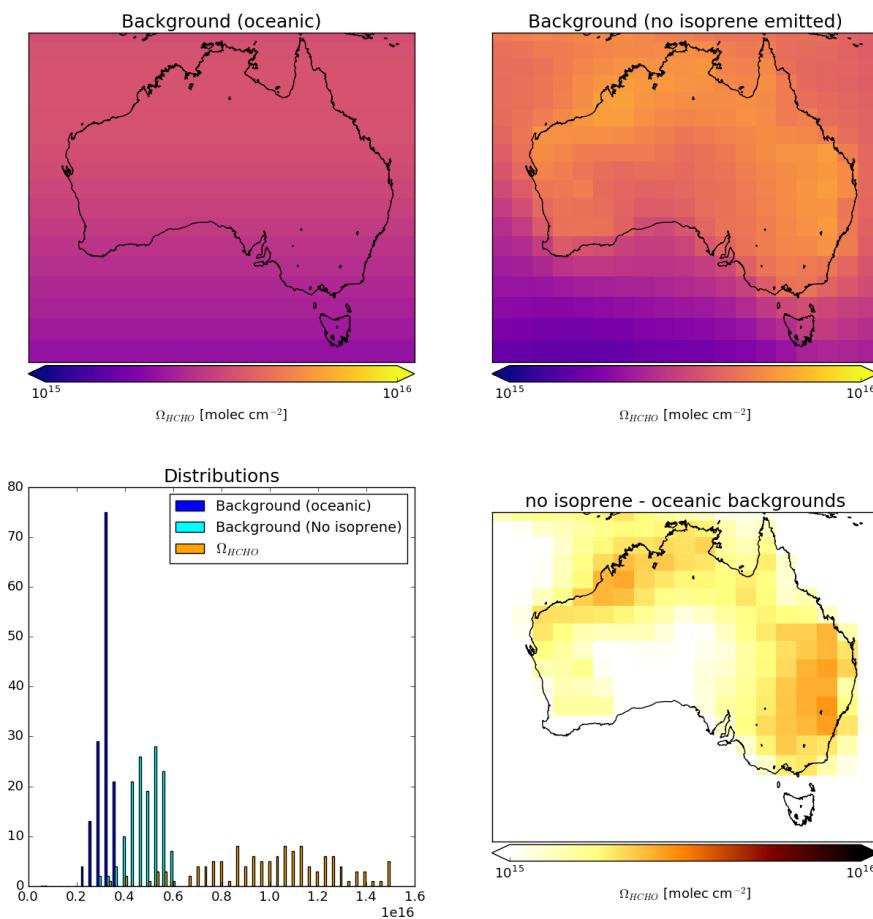


FIGURE 3.15: Top left: Background total column HCHO based on longitudinally averaged values over the remote Pacific Ocean. Top right: Background total column HCHO based on GEOS-Chem output with isoprene emissions set to zero. Bottom left: Distributions of column amounts over land squares in Australia from the two background definitions and the monthly HCHO average from GEOS-Chem (standard tropchem run). Bottom right: Absolute difference between the two background defined value maps.

These negative calculated emissions are set to zero, as negative emissions physically meaningless in this calculation. Relative uncertainty in these grid squares is set to 100% for later estimations of uncertainty (Section 3.4.1).

### 3.2.9 Running GEOS-Chem using a posteriori emissions

After creating the a posteriori isoprene emissions estimate (at monthly resolution), GEOS-Chem is re-run with biogenic emissions scaled to match the new estimate. This is performed by applying a seasonal scaling factor  $\alpha$ , based on the multi-year monthly average difference between midday a priori and a posteriori emissions at  $2^\circ \times 2.5^\circ$  horizontal resolution.  $\alpha$  is the ratio between the multi-year averaged monthly emissions from GEOS-Chem  $E_{GC}$  and the a posteriori  $E_{OMI}$ :

$$\alpha = \frac{E_{OMI}}{E_{GC}} \quad (3.12)$$

This seasonal scaling retains shorter time-scale variability and meteorological dependencies built into the parameterisations of the MEGAN model, while ensuring the multi-year monthly averaged emissions match the a posterior totals. This method provides a preliminary estimate of the effects of scaling isoprene emissions towards the satellite based estimate, and running this analysis at a finer temporal resolution could be performed in future work. Initially  $\alpha$  is uniformly set to 1 globally. Where top-down emissions exist and  $E_{GC}$  is non-zero,  $\alpha$  is set using Equation 3.12.  $\alpha$  is applied through the emissions module in GEOS-Chem where isoprene emissions are calculated. First, the new midday (13:00-14:00 LT) emissions (per grid box) are combined forming a multi-year monthly mean, which can be compared to the a priori equivalent. Missing values for  $\alpha$  when  $E_{GC}$  are zero are a negligible issue since the dominant discrepancies between estimates occur during summer when high emission rates are overestimated.

To create the  $\alpha$  scaling factor, modelled and top-down emissions are compared, which requires a change of units. Top-down emissions calculated in this work are in units of atom C  $\text{cm}^{-2} \text{ s}^{-1}$ , while modelled emissions are in  $\text{kg cm}^{-2} \text{ s}^{-1}$ . In order to calculate the top-down emissions in kg, each grid square is multiplied by its area, and then daily emissions are assumed to follow a sine wave peaking at the mid-day value. The sine wave approximates a daily insolation amplitude, allowing one to convert peak emission rates into daily emission amounts through integration. Figure 3.16 shows how the daily approximation of total emitted isoprene per grid square is calculated. Daytime hours are estimated per month, from 14 hrs (Jan) to 10 hr (Jul) (<https://en.wikipedia.org/wiki/Daytime>). This approximation is required since OMI observations occur at midday, when isoprene emissions are at their diurnal peak. GEOS-Chem emissions are similarly multiplied by area, but then integrated over time using hourly output to derive emissions in kg.

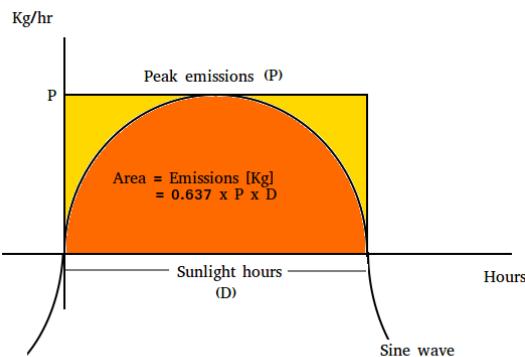


FIGURE 3.16: Total daily isoprene emissions (in kg) is represented by the area under the sine wave.

### 3.3 Results

Australia covers roughly  $7.7 \times 10^6 \text{ km}^2$ , with heterogeneous environmental conditions. The results presented in this section are therefore frequently split into five regions that are differentiated by colour, as shown in Figure 3.17. These regions are large enough to reduce the uncertainty with at least 10 grid squares in each area, and small enough to be somewhat homogeneous at the continental scale. Oceanic grid squares are filtered out of regional averages that follow.

#### 3.3.1 A posteriori emissions

Figure 3.18 shows a priori emissions over Australia along with a posteriori emissions calculated as described in the prior sections. This figure shows the time series of seasonal area averaged midday emissions, and their absolute differences. Differences between a priori and a posteriori estimates are seasonally and spatially diverse. A seasonal overestimate peaking in summer can be seen in all regions except for the northern region, which is generally overestimated but less so in summer. [Figure 3.19 shows how the yearly decrease is distributed spatially over Australia, with  \$E\_{GC}\$  and  \$E\_{OMI}\$  in  \$\text{Tg yr}^{-1}\$  calculated as a multi-year mean. Across all of Australia large reductions of total emissions are seen using the new top-down estimate.<sup>tb</sup>](#)

Figure 3.20 shows the multi-year seasonal emissions for each region for the a priori and a posteriori emissions side by side. The a priori is approximately twice that of the a posteriori. Absolute differences are highest in spring and summer, when emissions are generally greatest. The exception is the northern region, where the largest overestimates occur in spring, and large differences are seen in all seasons. Uncertainty is summarised for the a posteriori using the mean monthly uncertainty per grid square (calculated as described in 3.4.1) within each region, divided by the square root of 24 (8 years times 3 months per season). The uncertainty shown for each region is the average over the grid squares within that region, without reduction by the square root of the number of grid squares. Uncertainty is relatively small since 8 years of data are combined into the seasonal mean. The potential bias (see Section 3.4.1) is added

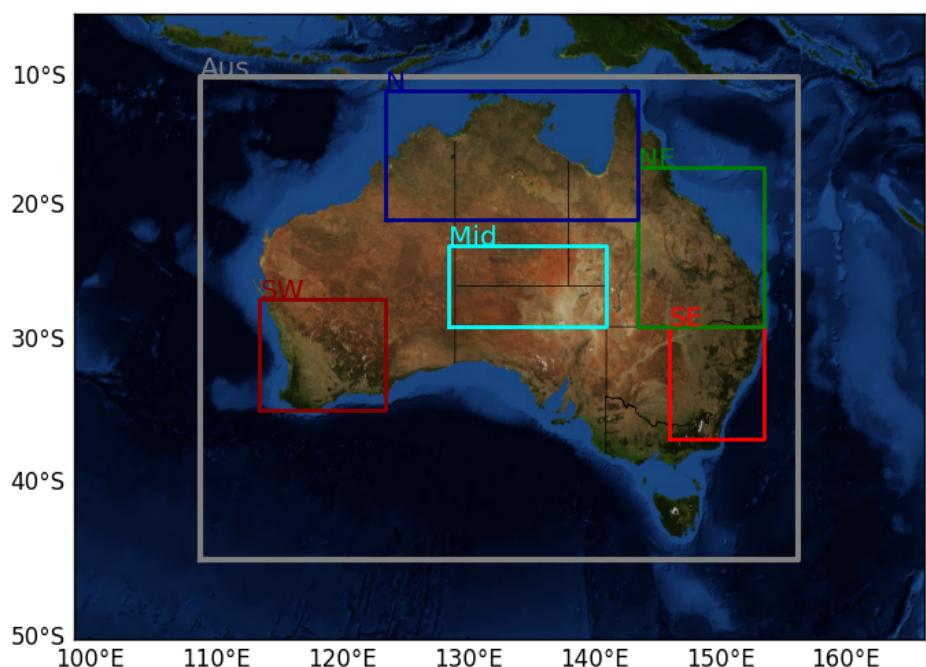


FIGURE 3.17: Sub-regions used in subsequent figures: Northern, North Eastern, South Eastern, South Western, and Middle. Australia-wide averages will be black or grey, while averages from within the coloured rectangles will match the colour shown here. Oceanic grid squares are filtered out from any area averages made in this chapter.

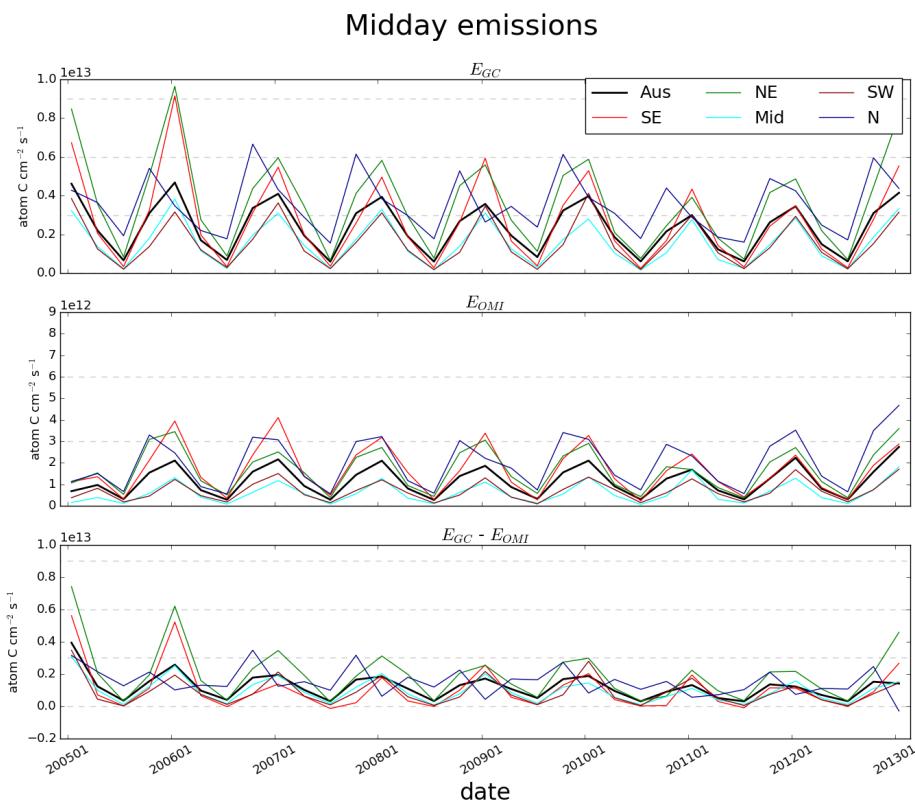


FIGURE 3.18: Row 1: Biogenic emissions of isoprene from GEOS-Chem (a priori,  $E_{GC}$ ). Row 2: Emissions calculated using the OMI top-down inversion (a posteriori,  $E_{OMI}$ ). Row 3: Absolute differences between the first two rows. Midday emissions are averaged for each season (DJF, MAM, JJA, SON), and colours represent averaged areas from subregions shown in Figure 3.17. Grey dashed horizontal bars are added to highlight the scale differences between rows.

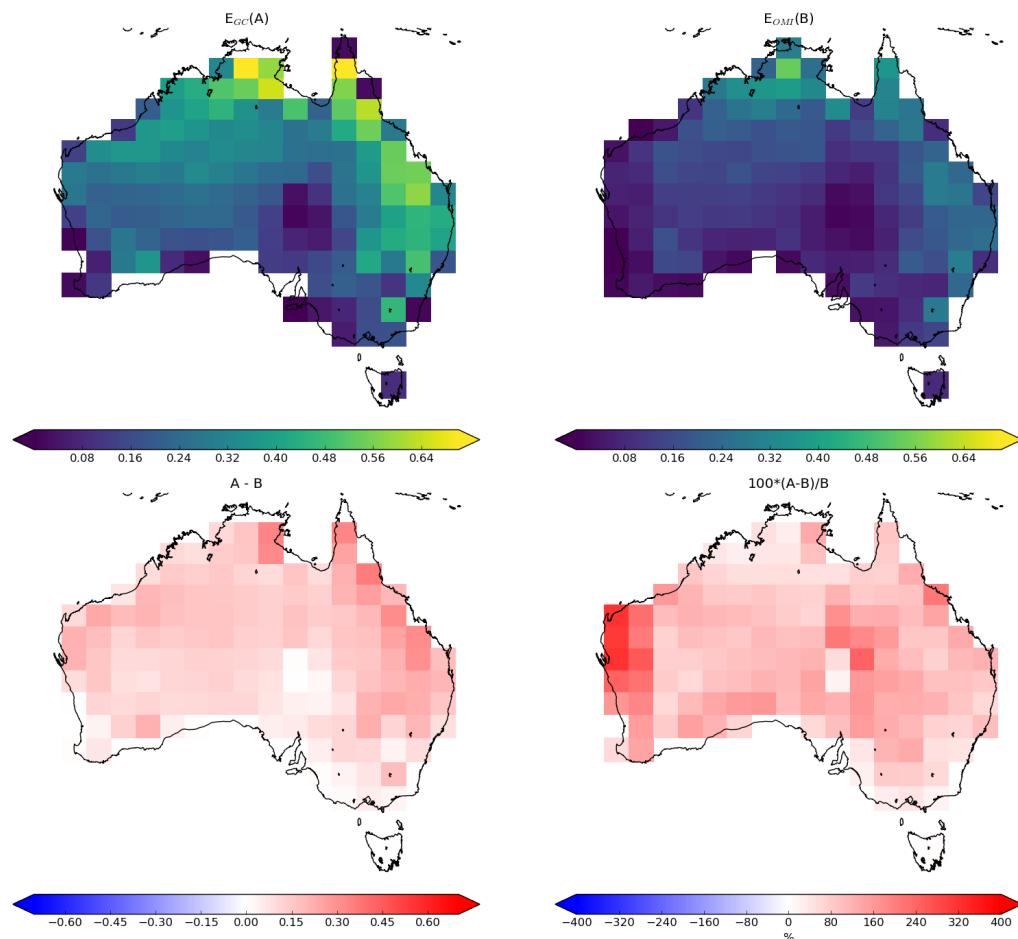


FIGURE 3.19: Top row: multi-year mean a priori emissions in  $\text{Tg yr}^{-1}$  from  $E_{GC}$  (GEOS-Chem; running MEGAN) and from  $E_{OMI}$  (top-down emissions) respectively. The total flux from  $E_{OMI}$  is calculated assuming a sinusoidal daily cycle, with daylight hours prescribed for each month as described in Section 3.2.9). Bottom left and right show the absolute and relative differences, respectively.

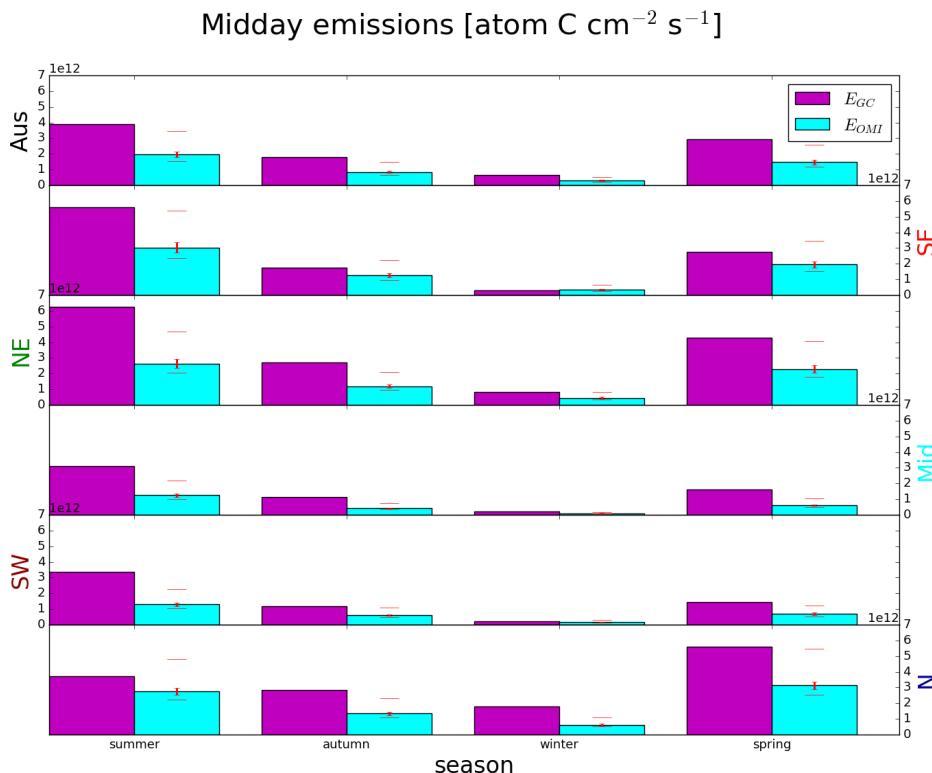


FIGURE 3.20: Regional multi-year seasonal mean a priori emissions (magenta) compared to a posteriori emissions (cyan). Error bars show the regionally averaged uncertainty (calculated as described in 3.4.1). Additional horizontal dashes show the uncertainty plus effects from potential HCHO biases (discussed in Section 3.4.1) from satellite underestimation (40%) and monthly clear sky overestimation (13%).

to the uncertainty and displayed using horizontal bars. The highest over-estimates (occurring in spring in the Northern region, and summer in all others) lie outside the potential bias caused by satellite underestimates of HCHO (Section 3.4.1). However, many overestimated seasons within each region are within this wide range of potential bias and uncertainty. Some monthly grid squares are very uncertain ( $> 200\%$ ) and these are removed for this plot, causing an increase in a posteriori emissions of  $< 1\%$ . There are very few of these grid squares, and they appear to be grid squares with very low emission rates and low pixel counts. The cause for these highly uncertain grid squares is not analysed further, but may be worth examining in future work.

Figure 3.21 shows the multi-year monthly mean and inter-quartile range of mid-day isoprene emissions estimates in each region. Months from September to April show the most difference between a posteriori and a priori. The most overlap is seen in the south-eastern region, where high summer emissions along with high variance occur in both the a priori and a posteriori. The highest variance is seen in both eastern regions, potentially due to diversity within the regions which include high density cities, large forests, and rural areas. This overestimate by the a priori may be caused

by some mixture of overly high emission factors and high emission sensitivity to temperatures and soil moisture (Emmerson et al. 2016; Emmerson et al. 2019). Over the entirety of Australia the seasonal cycle of emissions is shown to be overestimated by the *a priori*, and further analysis and measurements are required to determine the cause.

While most regions show similar overestimates, the northern region of Australia follows a different cycle of bias. Northern Australian emissions appear to be overestimated throughout the year, with the smallest bias in early summer. One potential reason is that during the wet season (November-April) there are fewer satellite measurements due to increased cloud coverage, along with a different ecosystem response to sunlight and temperature (e.g., Surl, Palmer, and Abad 2018). This is evidenced by the approximately 20% lower summer OMI pixel count (before filtering) in both northern regions (see Section 3.4.3). Low measurement counts in summer in the northern region could lead to a low bias in the *a posteriori* emissions estimate from the drier regions associated with lower biogenic emissions being over-represented. In situ measurements in both monsoon and non-monsoon seasons are required before robust conclusions can be drawn in this region.

There is only weak correlation between daily estimates of the *a priori* and *a posteriori* ( $r < 0.2$ ), and *a priori* distributions show more variance (figure in Appendix B.1). This is likely due to the filtering applied to satellite data (e.g., whenever cloud coverage exceeds 40%) which reduces the count and spread of *a posteriori* emission calculations. Figure 3.22 shows how the distributions of *a posteriori* emissions compare to *a priori* emissions in each region during summer months (December - February) with zeros removed from both distributions. This figure also shows the regressions between monthly averages of the same data. In the summer monthly averages, the linear regression coefficient  $r$  ranges from 0.2 to 0.81 depending on which region is being compared. The highest correlations between *a priori* and *a posteriori* emissions are in the southeast ( $r = 0.81$ ) and southwest ( $r = 0.79$ ) regions, followed by the northeast ( $r = 0.61$ ) region. This is likely due to isoprene emissions in these regions being dominated by the biogenic sources (large forests) that the top-down emission estimate is based upon. Although the northern region also contains large areas of forest, the correlation is the worst. In part this is due to misrepresented forest emissions in the model, exacerbated by unrepresented responses to moisture and poorly modelled environmental stresses in this region which lies within the tropics and undergoes monsoonal weather and intense heat and drought seasons (e.g., Emmerson et al. 2016; Surl, Palmer, and Abad 2018; Emmerson et al. 2019). If the same analysis is performed over winter months, the regression in the northern region is greatly improved (to  $r = 0.66$ ), while other regions remain approximately the same. If the modelled emissions distributions are correct (even if the absolute magnitudes are not), this suggests that satellite measurements are not capturing representative samples of monthly grid square averages in the north in summer, likely because of cloud coverage, and the summer inversion in this region may be biased.

When comparing total *a priori* emissions summed over Australia to the *a posteriori* calculated using the top-down inversion, a decrease of  $\sim 46\%$  from  $39 \text{ Tg yr}^{-1}$  to  $21 \text{ Tg yr}^{-1}$  is apparent. Table 3.5 compares annual Australian isoprene emissions estimated in this work to previously published values. [The \*a posteriori\* estimated here](#)

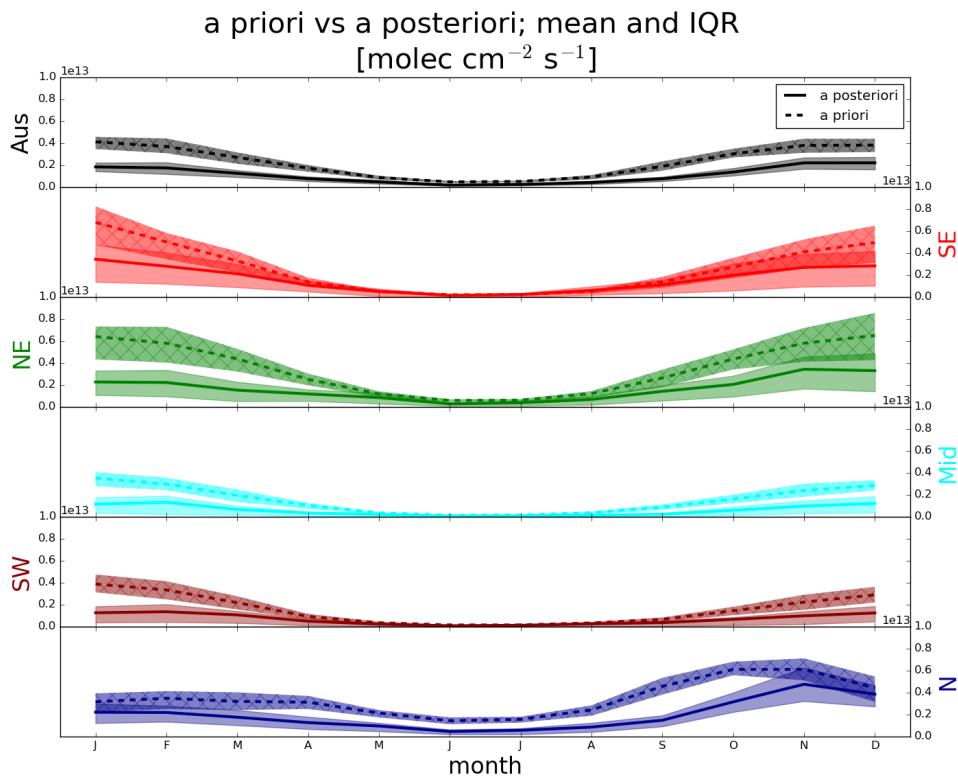


FIGURE 3.21: The multi-year monthly mean (lines) and inter-quartile-range (shaded) of midday (13:00-14:00 LT) isoprene emissions estimates. A priori emissions are shown by the dashed lines and hatched shaded areas show the inter-quartile-range. A posteriori emissions are shown using the solid lines, with inter-quartile-range shown by un-hatched shaded areas. Colours denote the region over which the multi-year monthly average was taken, as in Figure 3.17.

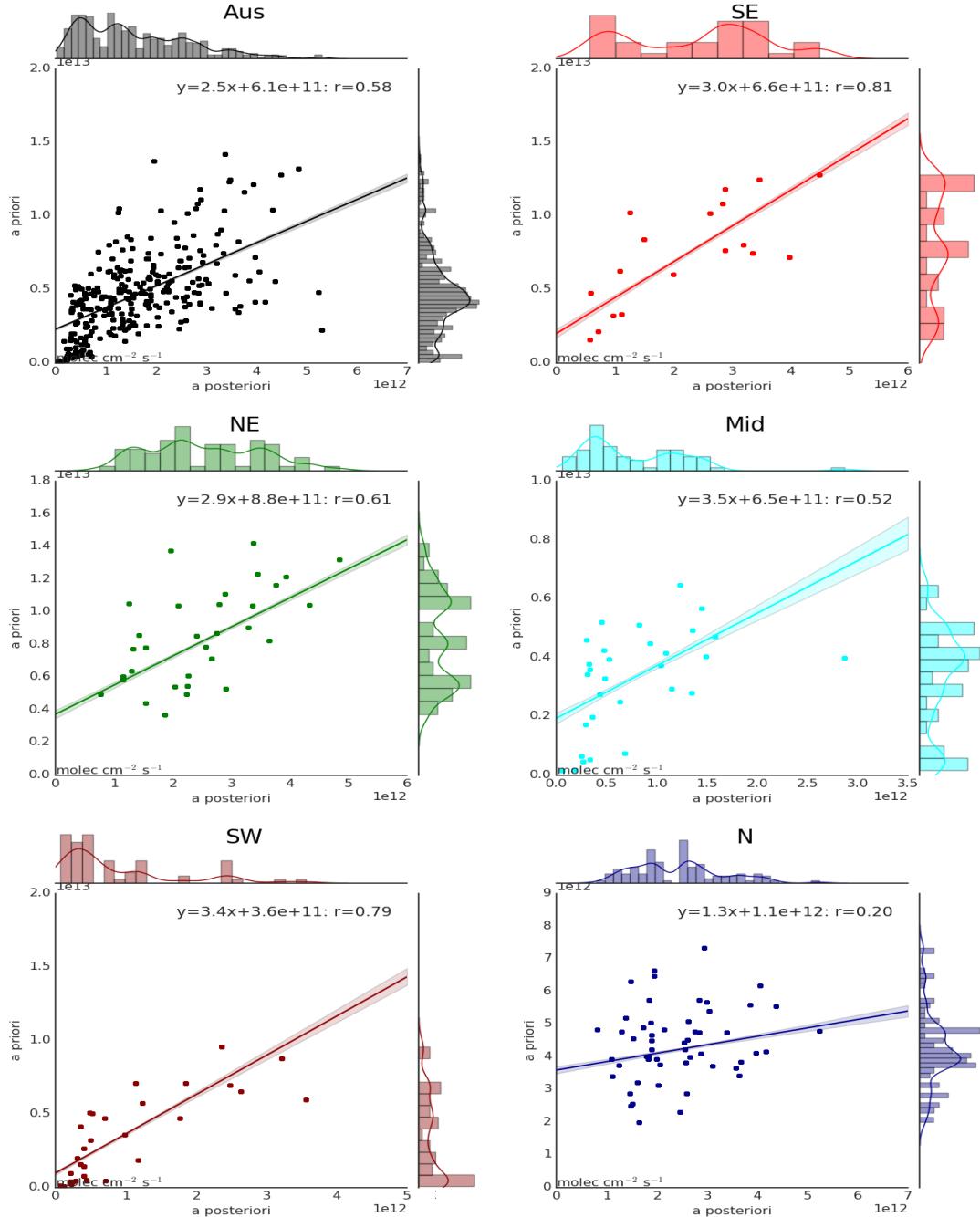


FIGURE 3.22: Scatter plot of a priori emissions against a posteriori emissions using monthly averaged grid squares as regression data points. Data points are created using monthly averages (of midday emissions) for each grid box for each month of summer (DJF) within each region shown. Multiple years of data are used, meaning if a region has 10 grid boxes, the 8 years of data will add up to  $10 \text{ boxes} \times 3 \text{ months} \times 8 \text{ years} = 240$  data points minus filtered and zero emission squares. Plots are coloured by regions shown in Figure 3.17. The linear best fit regression is inset into each plot along with the line equation and regression coefficient. The normalised distribution of each population is shown at the top and right spine of each subplot, with the right spine (facing the a priori axis) using the a priori axis and scale, and the top spine using the a posteriori axis and scale.

TABLE 3.5: Isoprene emissions (Tg/yr) from Australia

Estimate <sup>a</sup>	Source	Year	Reference
39.2(4.0)	bottom-up	2005-2012	This thesis <sup>b</sup>
~ 80	bottom-up	1980-2010	Sindelarova et al. (2014) <sup>c</sup>
26-94	bottom-up	2005-2013	Bauwens et al. (2016) <sup>d</sup>
20.7(1.6)	top-down	2005-2012	This thesis <sup>e</sup>
36	top-down	2005-2013	Bauwens et al. (2016) <sup>f</sup>

a: Interannual standard deviation shown in parentheses.

b: GEOS-Chem with MEGAN diagnostics based on 3-hourly averages.

c: MEGAN run using MERRA meteorology.

d: Range shown here based on three models, two of which implement MEGAN.

e: Based on daily peak emissions integrated over a sinusoidal daily curve.

f: OMI based top-down inversion.

to be  $21 \text{ Tg yr}^{-1}$ . Our a posteriori estimate of  $21 \text{ Tg yr}^{-1}$ <sup>mb</sup> suggests isoprene emissions may be lower than all prior bottom up estimates. This, and is close to the lowest bottom-up~~top-down~~ estimate of Bauwens et al. (2016) of  $26 \text{ Tg yr}^{-1}$ . There are several major differences between work performed in this thesis and work done by Bauwens et al. (2016). They use a fully adjoint version of the IMAGESv2 CTM with emissions determined by MEGAN-MOHYCAN, and iteratively update biogenic isoprene emissions and pyrogenic VOC emissions so that the CTM produces HCHO columns that match (as near as possible) the column HCHO observed from OMI. The differences in results between their work and work performed here are most likely to be primarily driven by differences in modelled chemistry, rather than differences in top-down methodology. Both techniques have the end result of reducing the differences between (respective) modelled and OMI total column HCHO. The fact that both top-down techniques suggest a priori bottom-up emissions are too high is a good indicator that these inventories are indeed overestimating isoprene emissions.<sup>tb</sup>

Figure 3.19 shows how this decrease is distributed spatially, with  $E_{GC}$  and  $E_{OMI}$  in  $\text{Tg yr}^{-1}$  calculated as a multi-year mean. Across all of Australia large reductions of total emissions are seen using the new top-down estimate.<sup>tb</sup>

### 3.3.1.1 Diurnal variability

Figure 3.23 shows the a priori daily emissions cycle for Australia compared to the estimated a posteriori emissions cycle over Australia. The conversion of midday a posteriori emissions ( $\text{ molec cm}^{-2} \text{ s}^{-1}$ ) into  $\text{Tg yr}^{-1}$  involves integration over an assumed sinusoidal diurnal emission cycle as described in Section 3.2.9. A priori emissions peak from approximately 11:00 LT to 16:00 LT, while outside these hours there is a non-sinusoidal drop in emissions to below the assumed a posteriori diurnal emission cycle. This means the conversion may be biased by this consistent difference between modelled a priori diurnal emission cycles and the assumed diurnal a posteriori cycle.

TABLE 3.6: 2005-2012 trend<sup>a</sup> in surface isoprene mixing ratio (ppbvC yr<sup>-1</sup>) from simulations using a priori and a posteriori isoprene emissions.

Region	a priori	a posteriori
Aus	<b>-.04</b>	<b>-.02</b>
SE	<b>-.15</b>	<b>-.07</b>
NE	<b>-.13</b>	<b>-.04</b>
Mid	<b>-.07</b>	<b>-.02</b>
SW	-.01	-.01
N	<b>-.06</b>	<b>-.02</b>

a: Statistically significant (two sided test with  $\alpha = 0.1$ ) trends are bolded.

This potential bias is not analysed further, and should be relatively small compared to other uncertainties as it only affects emissions towards the daily minima.

### 3.3.1.2 Trends

Figure 3.24 shows monthly deseasonalised a priori and a posteriori midday emission anomalies for each region. First the emissions are spatially averaged within each region to form a daily time series of midday emission rates. These are averaged into monthly data, and then the multi-year monthly mean is subtracted to form the anomaly time series. Any anomaly greater than three standard deviations from the mean is removed (crosses in Figure 3.24). An ordinary least squares linear regression is then performed to look for any significant trend. A trend is considered significant if the p-value from a Wald test (equivalent to a t-test) is less than 0.05. Trend results for isoprene are summarised in Table 3.6. The a priori midday surface isoprene concentrations show a small decline outside of the southwest region over the 8 year period from 2005-2012. This decline reduces in scale by approximately a factor of 2-3 in the a posteriori emissions. There is no significant change in trend due to scaling isoprene emissions. Since the scaling factor  $\alpha$  is applied to each grid square seasonally and not changing each year, changing trends are not expected.

### 3.3.2 Modelled impacts of reduced isoprene emissions

This section uses GEOS-Chem to determine how the improvements to biogenic isoprene emissions impact subsequent atmospheric chemistry and composition. A posteriori emissions are implemented in GEOS-Chem as described in Section 3.2.9. Outputs from the scaled GEOS-Chem run using the a posteriori are denoted by superscript  $\alpha$ . For example, column HCHO from GEOS-Chem before and after scaling are denoted  $\Omega_{GC}$  and  $\Omega_{GC}^\alpha$  respectively. Figure 3.25 shows  $\alpha$  for January and June averaged over 2005-2012, along with the time series of  $E_{GC}$  and  $E_{OMI}$  and  $\alpha$  calculated for the 2°x 2.5° grid box containing Sydney and their multi-year seasonal average. GEOS-Chem

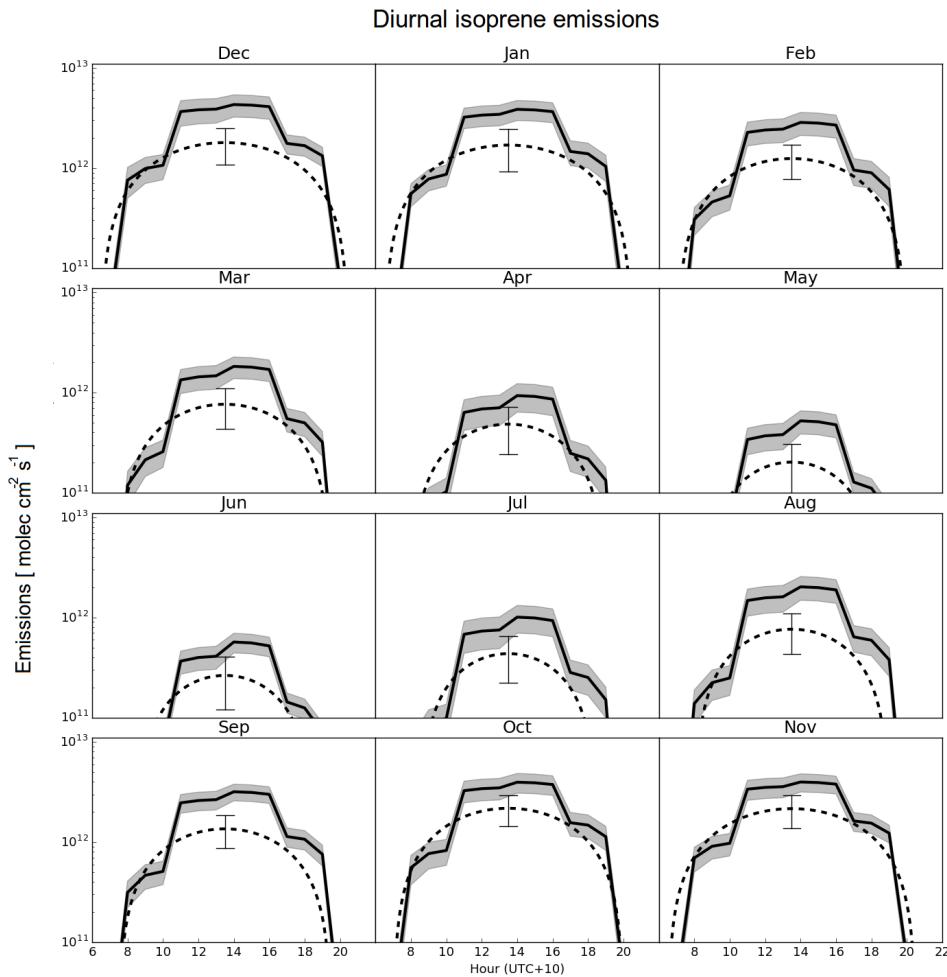


FIGURE 3.23: The diurnal cycle of GEOS-Chem a priori emissions (solid line) averaged by month into hourly bins over from 2005 to 2013 along with top-down a posteriori (dashed line) emissions. Standard deviations for the monthly average are shaded for the a priori, and shown with error bars at 13:30 LT for the a posteriori. Top-down emissions shown here are based on monthly midday emissions being the peak of a sine wave which drops to zero after and before daylight hours (see Section 3.2.9).

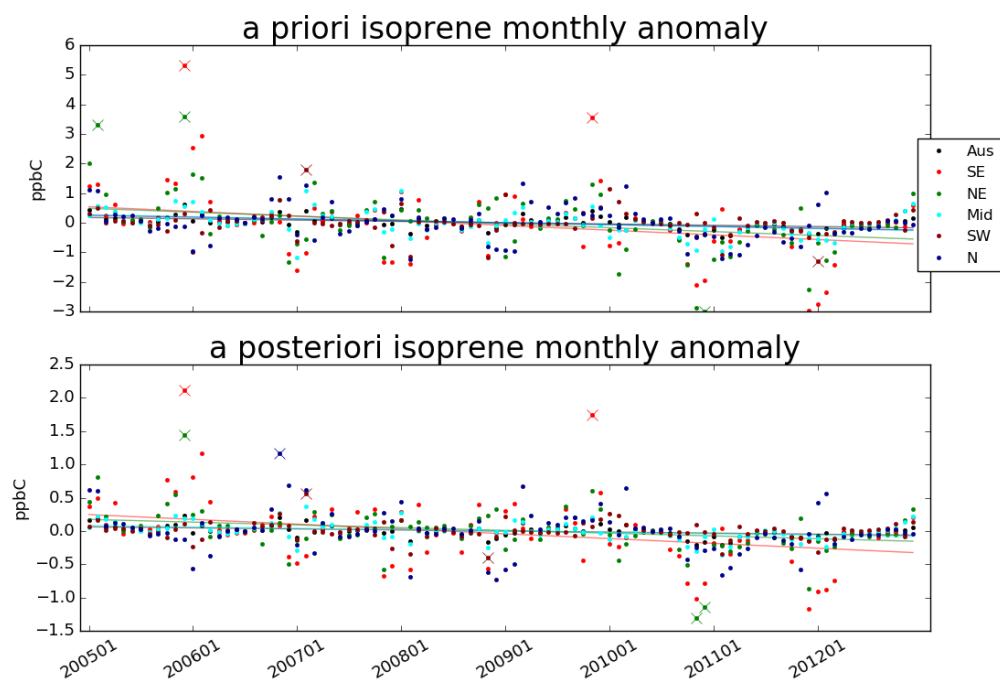


FIGURE 3.24: A priori (row 1) and a posteriori (row 2) monthly emissions anomalies from multi-year monthly mean, split by region (see Figure 3.17).

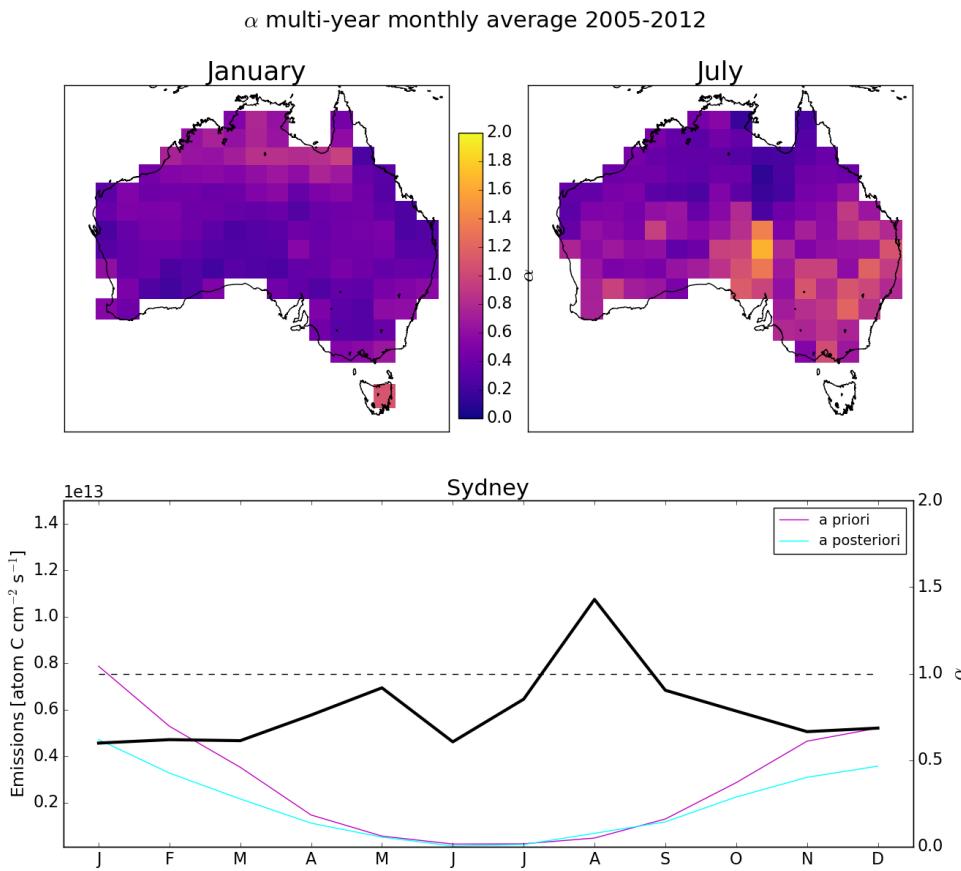


FIGURE 3.25: Row 1: Isoprene emissions scaling factor  $\alpha$  for the average January (left) and July (right) over 2005-2012. Row 2: a priori (magenta, left axis), a posteriori (cyan, left axis), and  $\alpha$  (black, right axis) multi-year monthly averages calculated for the  $2^\circ \times 2.5^\circ$  grid box containing Sydney.

emissions are scaled according to this multi-year averaged gridded  $\alpha$ , which is seasonal and spatially diverse.

### 3.3.2.1 Implications for HCHO

As a preliminary check on the GEOS-Chem output, simulated  $\Omega_{GC}$  and  $\Omega_{GC}^\alpha$  are compared to  $\Omega_{OMI}$  over January and February, 2005 in Figure 3.26. In every region,  $\Omega_{GC}^\alpha$  is closer to  $\Omega_{OMI}$  with biases decreasing from  $\sim 100\%$  to  $\sim 50\%$  everywhere except the northern region, which has biases decreasing from  $\sim 50\%$  to  $\sim 25\%$ . Note that this is not an independent validation as  $\Omega_{OMI}$  drive the creation of  $\Omega_{GC}^\alpha$ , and as is expected the relationship is much improved. The remaining differences are most likely driven by filtering and temporal averaging of the applied scaling factor  $\alpha$ . When looking over all of Australia from 2005-2012, the summer mean decreases from  $9.8 \times 10^{15}$  to

$7.4 \times 10^{15}$  molec cm $^{-2}$  after scaling isoprene emissions, while the satellite column is  $4.9 \times 10^{15}$ .

Figure 3.27 shows HCHO is most reduced in summer for the majority of Australia. In winter, a reduction is seen along the northern coast of Australia, most likely because emissions in this area are less affected by the seasonal decline. Seasonal means and the standard deviations (between all grid boxes and days used in averaging) for each region are summarised in Figure 3.28. Both mean and variance are reduced after running GEOS-Chem with scaled isoprene, although the mean is lower still in OMI vertical columns. Model output standard deviations in summer range from  $\sim 20 - 30\%$ , while OMI standard deviations range from  $\sim 32 - 41\%$ . The highest OMI standard deviations occur in winter, ranging over  $\sim 46 - 73\%$ , which is the opposite of model output with standard deviations between  $\sim 12 - 27\%$ . This could be an effect of the increased winter uncertainty in satellite output, which is amplified by the low column amounts in the season. Overall the standard deviation within model output appears to be too low in all regions, ranging from three quarters (south eastern region) to one fifth (middle region) of the standard deviations within the OMI HCHO column output. These standard deviations are formed from the monthly averaged grid boxes used to create each multi-year seasonal mean. Their low value (relative to satellite measurements) suggests that the model does not capture the full range of variance over time and space of HCHO levels over Australia; however, variance is heightened in satellite measurements due to the filtering that reduces the number of grid squares that form the multi-year seasonal mean.

Decreasing isoprene emissions in the model lead to reduction in HCHO concentrations as one would expect. Figure 3.29 shows an example (over Wollongong) of the modelled HCHO profile before and after scaling isoprene emissions. This reduction of HCHO is greatest at the surface, and is highly correlated ( $r > .8$ ) in all regions to the reduction in isoprene emissions. The high correlation is effectively a sanity check of the top-down method used in this thesis to estimate isoprene emissions.

### 3.3.2.2 Implications for ozone

Isoprene oxidation can eventually lead to ozone formation, especially when isoprene-enriched air masses mix with polluted urban air masses that contain high NO<sub>x</sub>. Figure 3.30 shows surface level (up to  $\sim 150$  m altitude) ozone concentrations over 2005 before and after scaling modelled isoprene emissions. Reducing isoprene emissions lowers surface ozone concentrations by  $\sim 1$  ppbv in all regions in all seasons. The largest reductions ( $\sim 3$  ppbv) occur in the northern region in spring. A regression between the change in isoprene emissions, and the change in surface ozone was performed (as in the prior section with HCHO); however, only very weak correlations between reductions are apparent. While the overall decrease in surface ozone is clear, there is no direct correlation between monthly grid square averaged reductions in isoprene emission and surface ozone concentrations in the same grid square. This suggests that changes in isoprene emissions affect ozone in non-local grid squares due to transport, including grid squares where absolute emission reductions are not as strong. [Figure 3.31 shows Australian summer surface level ozone with and without scaling isoprene emissions to match the multi-year top-down estimation. Surface ozone is reduced in](#)

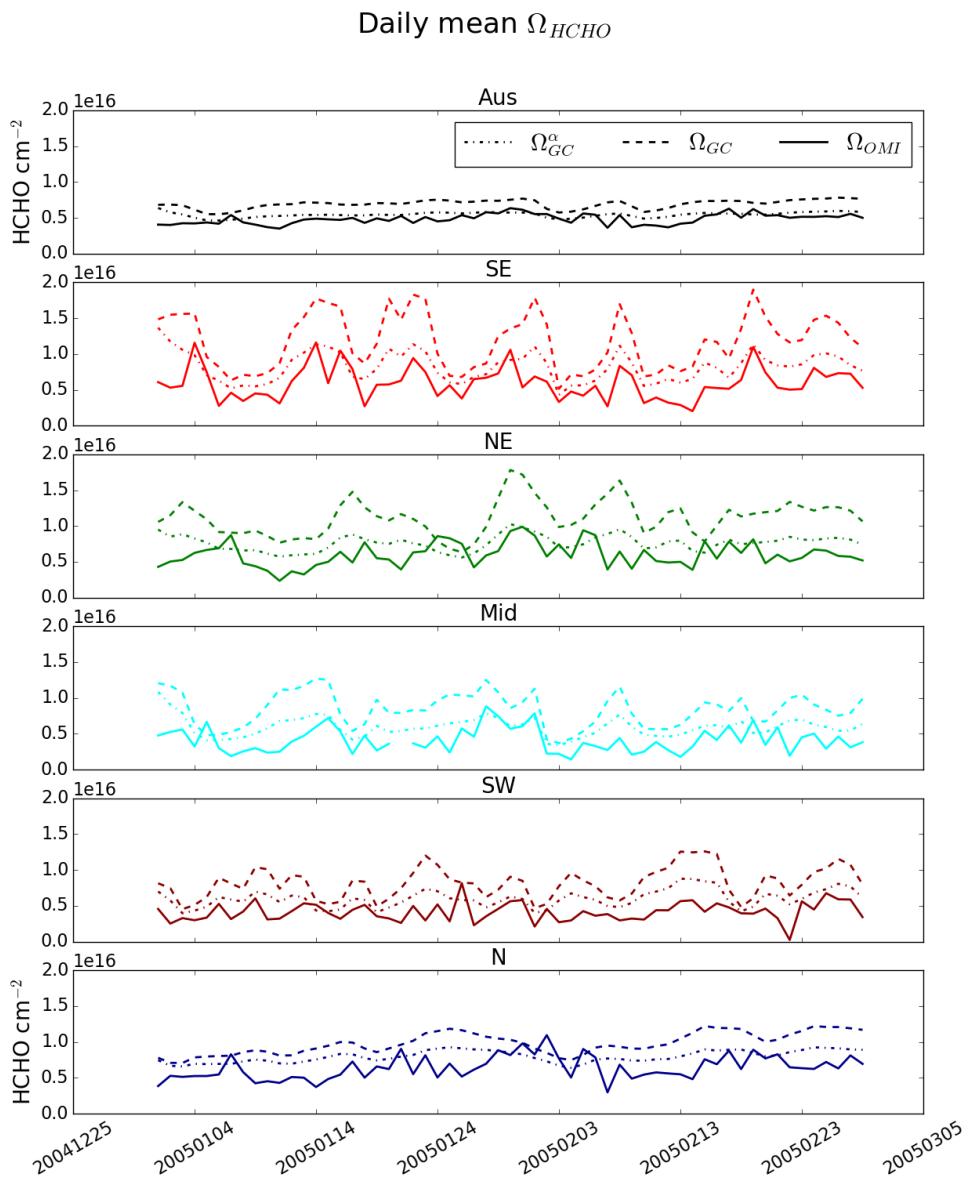


FIGURE 3.26: Daily mean total column HCHO amounts from GEOS-Chem with a priori (new emissions run) and a posteriori (tropchem run) a posteriori scaled isoprene emissions, along with the recalculated OMI HCHO columns. Each row shows the average over regions in Figure 3.17.

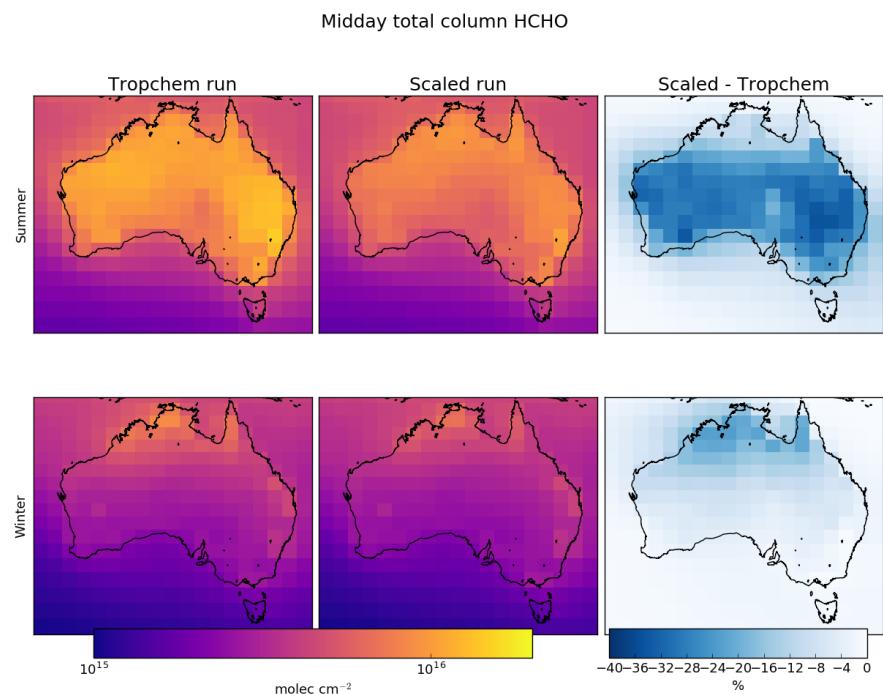


FIGURE 3.27: Total column HCHO a priori (left) and a posteriori (middle) isoprene emissions, and their relative differences (right). Top row shows summer (DJF) averaged total columns, while bottom row shows the winter (JJA).

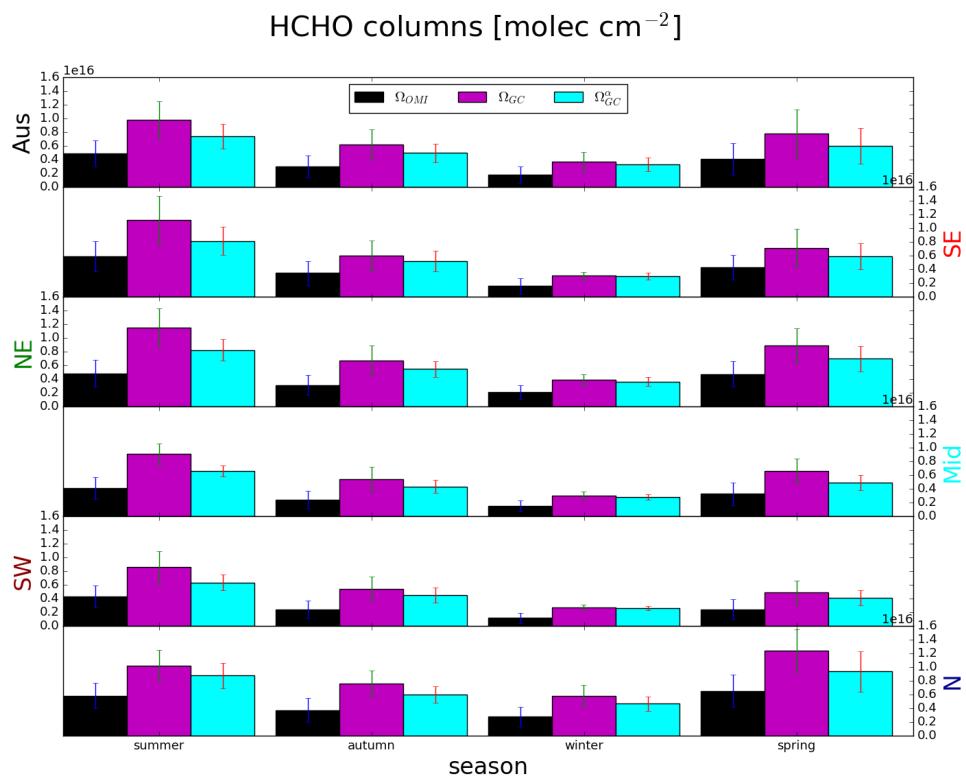


FIGURE 3.28: Regionally and seasonally averaged HCHO total columns from GEOS-Chem ( $\Omega_{GC}$ , and  $\Omega_{GC}^a$ ) and recalculated OMI measurements ( $\Omega_{OMI}$ ) side by side. Each row represents one region within Australia, while each column represents from left to right: summer, autumn, winter, spring. Standard deviations between grid boxes used to form the multi-year seasonal average are shown with error bars.

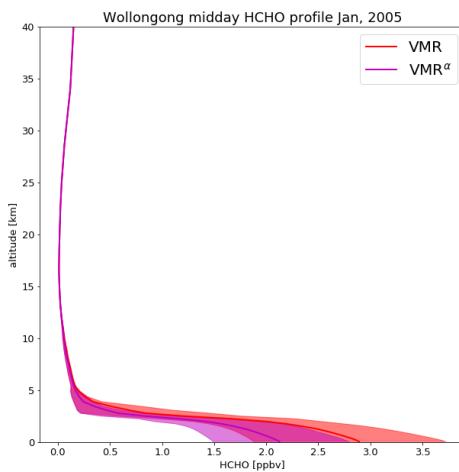


FIGURE 3.29: Monthly averaged HCHO profile over the  $2^\circ \times 2^\circ$  grid box containing Wollongong modelled by GEOS-Chem before (VMR) and after ( $\text{VMR}^\alpha$ ) scaling isoprene emissions. Shaded areas represent the inter-quartile range over the month.

summer by approximately 5%, with hotspots of reduction over southern New South Wales, Victoria, and Western Australia.<sup>tb</sup>

Ozone and fine particulate concentrations in Australian cities have not reduced over the last 10 years, unlike other atmospheric pollutants such as CO, NO<sub>2</sub>, and SO<sub>2</sub> (Keywood, Emmerson, and Hibberd 2016). This could be in part due to the downwind effects of isoprene emission, which are most likely to affect suburban fringes (e.g. western Sydney) of Australian cities which are surrounded by vegetation (Millet et al. 2016). Outside of densely populated regions, Australia is likely to be NO<sub>x</sub>-limited and changes in VOC emissions will have less of an impact on ozone production. Here the modelled ozone output is averaged over large areas that are mostly non-urban, which means detected ozone sensitivity to isoprene emissions is likely underestimated for cities. Figure 3.32 shows how ozone is reduced in summer and winter after scaling isoprene emissions. In summer, reductions are strongest in Sydney and Melbourne, but can also be seen in the west coast and central Australia. Winter reductions are more uncertain (relatively); however, they are strongest in the coastal and oceanic regions around Darwin. ~~In future work, the estimated emissions will be downscaled to better evaluate their impacts on ozone in different NO<sub>x</sub> regimes (for instance over cities).~~<sup>tb</sup>

Table 3.7 shows regionally how the changed isoprene emissions affected ozone concentration yearly averages. The overall reduction is  $\sim 3\%$ , with the largest reduction occurring in the north region. The strongest reductions in isoprene emission do not coincide with those of ozone concentration. This is likely due to both transport (ozone formation occurs downwind of isoprene emissions) and different NO<sub>x</sub> concentrations within each region. In future work, the estimated emissions will be

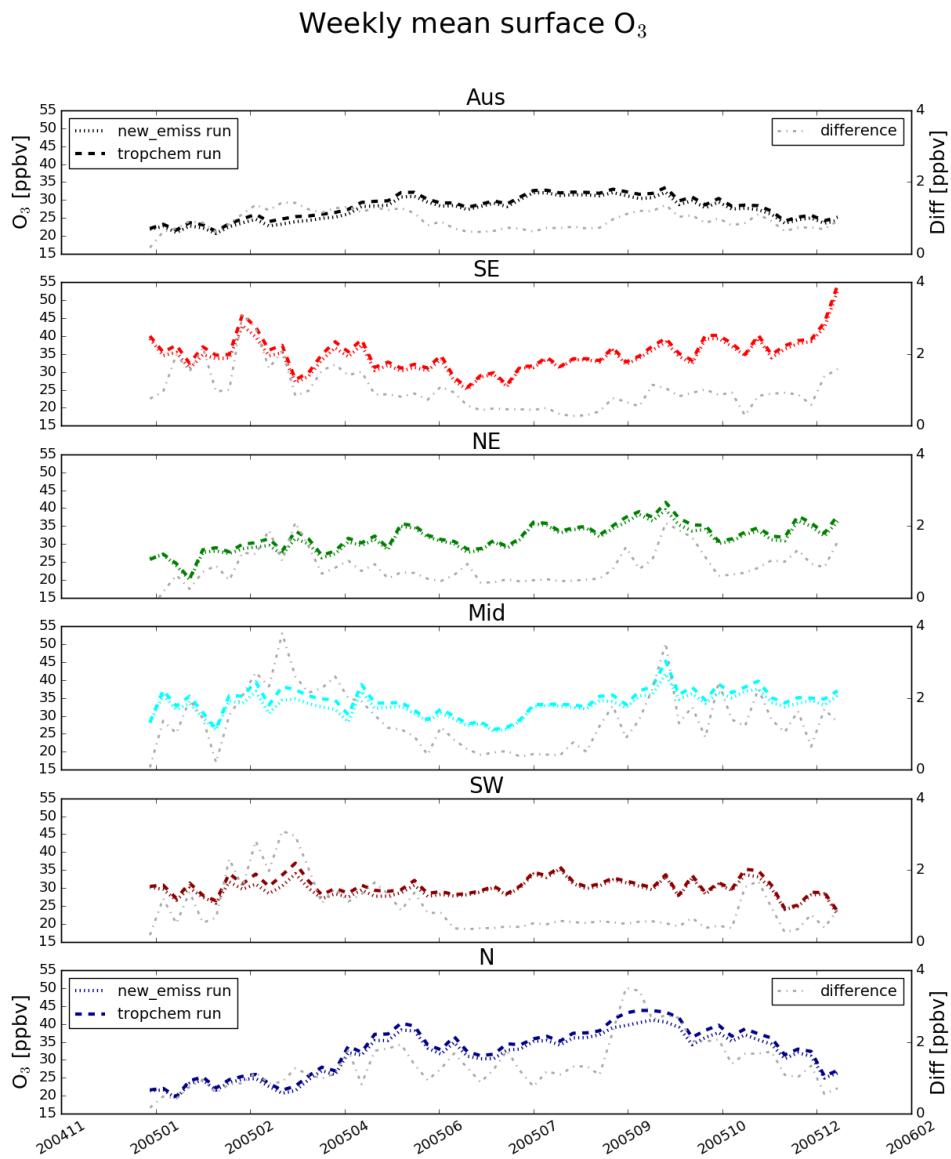


FIGURE 3.30: Surface ozone concentrations (ppb) per region over 2005. Concentrations are shown using the left axis, and absolute differences (tropchem run - scaled run) shown in grey using the right axis.

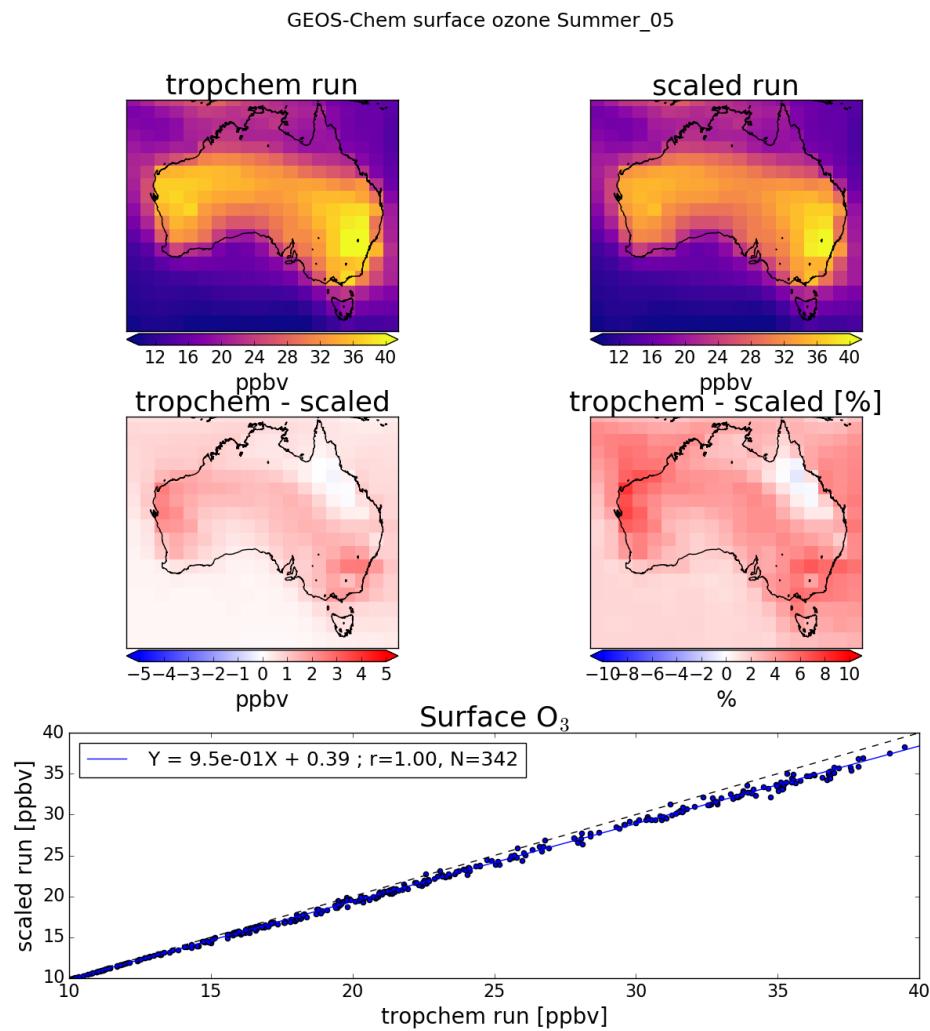


FIGURE 3.31: Top row: Summer surface-level ozone before (left) and after (right) scaling isoprene emissions in GEOS-Chem to match top-down estimates. Middle row: absolute (left) and relative (right) differences. Bottom row: linear regression between the two model runs along with a black dashed line representing the 1-1 ratio. Each point represents one summer averaged land grid square (as shown in the top row).<sup>tb</sup>

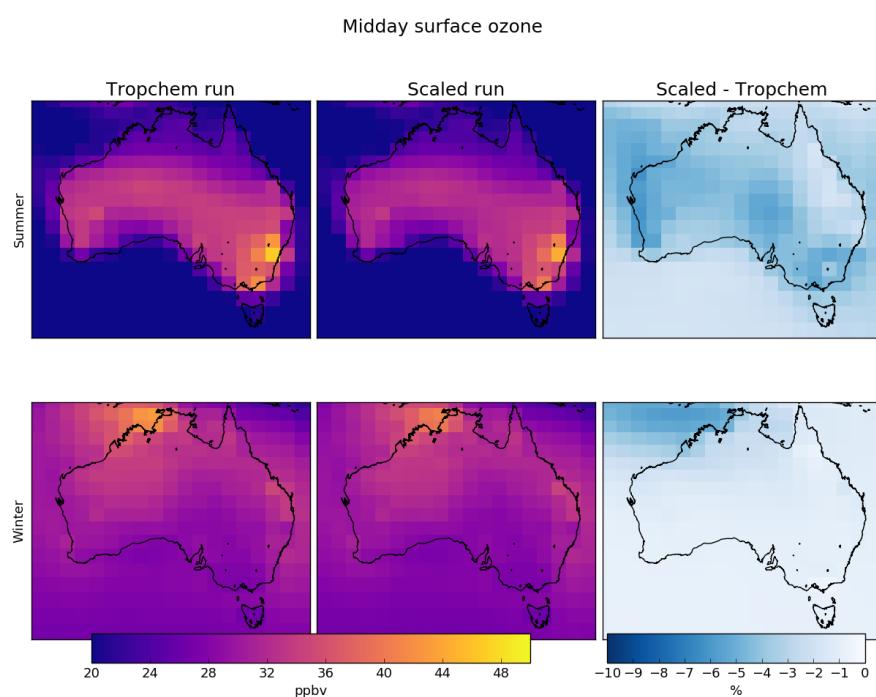


FIGURE 3.32: Multi-year seasonal surface (up to  $\sim 150$  m) ozone before (left) and after (middle) scaling isoprene emissions, and their relative differences (right). Top row shows summer (DJF) averaged total columns, while bottom row shows the winter (JJA).

TABLE 3.7: Isoprene emissions from MEGAN and top-down estimate in Tg C a<sup>-1</sup>, along with ozone surface amounts in ppb.

	AUS	SE	NE	MID	SW	N
<b>Bottom-up</b>						
Isoprene	34.67	2.80	6.33	2.31	2.04	9.95
Ozone	27.72	33.97	30.64	33.32	30.51	32.13
<b>Top-Down</b>						
Isoprene	18.26	1.61	3.02	1.03	0.99	6.31
Ozone <sup>a</sup>	26.93	33.10	29.99	32.19	29.69	30.92

a: Outputs from GEOS-Chem run with isoprene emissions scaled to match the top-down estimate.

downscaled to better evaluate their impacts on ozone in different NO<sub>x</sub> regimes (for instance over cities).<sup>b</sup>

### 3.3.3 Comparison with in situ measurements

Comparison between ground-based measurements and large ( $2^\circ \times 2.5^\circ$ ) averaged model output suffers from representation error. Figure 3.33 shows the SPS and MUMBA measurement campaigns (see Section 2.2), along with the extent of the relevant  $2^\circ \times 2.5^\circ$  GEOS-Chem grid box, a rectangle with edge lengths of roughly 200 km<sup>2</sup>. The urban footprint of Sydney and Wollongong can be seen, along with ocean, forest, and rural regions, which will all affect the model output as it is based on the average of inputs within the grid box. Due to high uncertainty in components of the top-down emissions estimate, temporal resolution is also limited. MUMBA (Section 2.2.3.1), SPS1 and SPS2 (Section 2.2.3.1) provide approximately one month (two for MUMBA) of hourly or daily data, which are compared in this section against surface level concentrations from GEOS-Chem (midday output) before and after scaling the biogenic emissions.

Figure 3.34 shows GEOS-Chem output in the grid square containing Sydney and Wollongong campaign measurements. Measurements between 13:00 and 14:00 are averaged daily. In order to minimise introduced bias, any measurements below the instrument detection limit are set to half of the detection limit, as performed in Lawson et al. (2015). Midday measurements from MUMBA (summer, Wollongong) compared against SPS (summer, autumn, Sydney) show lower isoprene and HCHO levels, and similar ozone levels, and also lower variance across all three species. Variation in SPS data are higher than modelled, which is likely due to a dependence on local meteorology, as plumes of HCHO or isoprene enriched air can be detected as they pass by while the modelled grid square averages these out. This higher variance is not seen in the MUMBA midday observations; however, this may be due to the small sample size of the measured data.

Coincident measured and modelled means and comparison statistics for each campaign for isoprene, HCHO, and ozone are summarised in Table 3.8. Isoprene correlations are greatest during the SPS1 campaign at  $r = 0.83, 0.84$ , nearly double those of SPS2 and MUMBA. The root mean square error (RMSE) in MUMBA is 1.7 ppb

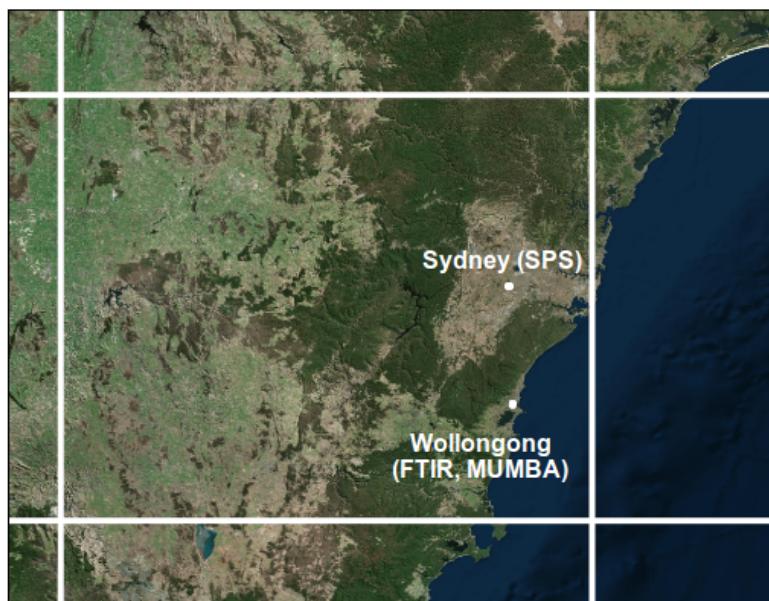


FIGURE 3.33: GEOS-Chem grid box ( $2^\circ \times 2.5^\circ$ ) containing Wollongong FTIR, SPS, and MUMBA campaign data.

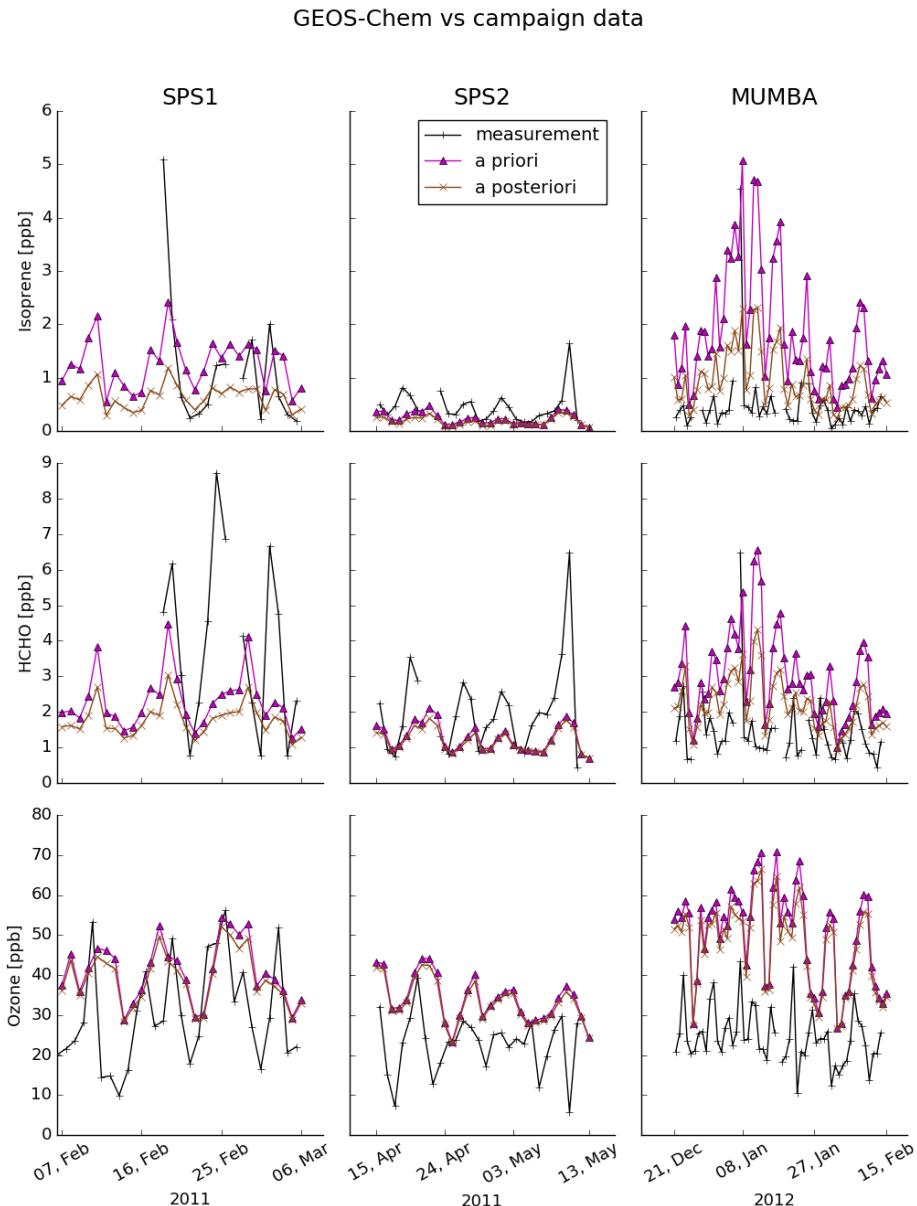


FIGURE 3.34: SPS1, SPS2, and MUMBA (left to right columns respectively) midday (13:00–14:00 local time) measurements of isoprene, HCHO, and ozone (top to bottom respectively). Shown in magenta and brown are the a priori and a posteriori GEOS-Chem surface outputs for the matching grid square at midday for days containing measurements.

TABLE 3.8: Campaign measurements compared to model output [ppb].

	SPS1			SPS2			MUMBA		
	mean	RMSE	r	mean	RMSE	r	mean	RMSE	r
Isoprene									
Meas	1.16			0.46			0.47		
GC	1.34	0.89	0.83	0.24	0.36	0.44	1.87	1.66	0.52
GC <sup>a</sup>	0.67	1.15	0.84	0.18	0.39	0.43	0.93	0.73	0.49
HCHO	mean	RMSE	r	mean	RMSE	r	mean	RMSE	r
Meas	3.92			2.02			1.41		
GC	2.38	2.63	0.47	1.28	1.28	0.77	3.00	1.97	0.44
GC <sup>a</sup>	1.83	3.01	0.54	1.19	1.35	0.78	2.21	1.17	0.47
Ozone	mean	RMSE	r	mean	RMSE	r	mean	RMSE	r
Meas	30.14			22.71			24.59		
GC	40.91	15.46	0.52	33.75	13.59	0.35	49.97	28.01	0.34
GC <sup>a</sup>	39.20	14.23	0.54	32.81	12.69	0.36	47.14	24.91	0.36

Meas: Measurements (midday means)

GC: GEOS-Chem (tropchem run)

GC<sup>a</sup>: (GEOS-Chem scaled run)

(~ 300%) and approximately 0.9 ppb and 0.4 ppb in SPS1, SPS2 respectively (~ 100%), before isoprene scaling is applied. In MUMBA the reduced isoprene emissions improves the model drastically, bringing the RMSE down to ~ 100%, however this improvement is not seen in either of SPS1 or SPS2. Similarly for HCHO, the MUMBA RMSE is reduced from 2.0 ppb (~ 130%) to 1.2 ppb (~ 80%) by bringing down the mean modelled amounts. Modelled HCHO is low compared against measurements from Sydney (SPS1 and SPS2), and is further lowered through scaling. Without improved model resolution or further measurements, this is assumed to be due to local influences seen in the measurements but not captured by the model, such as HCHO enriched air from local sources. Modelled ozone concentrations are slightly reduced (3 – 8%) when scaling isoprene emissions, however the concentrations remain 30 – 50% too high for Sydney and approximately 100% too high for Wollongong. This is likely due to the reduced ozone concentrations in cities compared to regional areas, as the modelled ozone concentrations are averages over the larger area which is mostly non-urban.

A spectrometer (FTIR) on the roof of the Chemistry building at the University of Wollongong measures total column HCHO in the atmosphere during clear sky conditions. This is the only non-satellite long-term measurement record of total column HCHO available in Australia (see Section 2.2.3.3). In order to compare modelled profiles against retrievals from the FTIR, modelled profiles are first convolved with the instrument averaging kernel and a priori. FTIR output is resampled to only include measurements taken at midday (13:00-14:00) and GEOS-Chem overpass outputs are interpolated onto matching vertical levels for days where FTIR output exists. Figure 3.35 shows total column HCHO from FTIR, and from GEOS-Chem before and after scaling isoprene emissions. While the comparison suffers from representational error between FTIR measurements and GEOS-Chem output (see Figure 3.33), one can see

TABLE 3.9: Mean total column HCHO amounts in  $10^{15}$  molec  $\text{cm}^{-2}$  at Wollongong.

Season	$\Omega_{FTIR}$	$\Omega_{GC}$	$\Omega_{GC}^{\alpha}$
summer	15.4	17.7	13.6
winter	3.68	4.41	4.49

that the summer overestimate of HCHO from GEOS-Chem is removed by isoprene scaling. The mean summer underestimate of HCHO shown after scaling makes more sense due to the relatively dense forested areas around Wollongong, which would raise local HCHO concentrations above the average for the large grid square represented by GEOS-Chem. Another feature is the January and December dip in all three datasets. The FTIR has a lower data count and higher variance in these (holiday) months as measurements involve manual processes, increasing uncertainty for the measurements. Analysis of this dip could be performed in future work; however, it would require an in depth examination of local and synoptic HCHO patterns, and higher resolution model data. Seasonal mean HCHO from the FTIR measurements and co-located, convolved, and resampled model output is summarised in Table 3.9. The original modelled summer vertical column is higher than the FTIR measurements by  $\sim 15\%$ , which decreases with isoprene scaling to  $\sim -12\%$ . In winter the column is modelled high by  $\sim 20\%$ , which increases to  $\sim 22\%$  after scaling.

### 3.4 Uncertainty

This section identifies and quantifies the overall uncertainties of calculating isoprene emissions using OMI HCHO observations and the GEOS-Chem model in the top-down method used in this chapter. However, these uncertainties lack verification against measurements. Even as the top-down inversion performed in this chapter attempts to work around the lack of measurements over Australia, it suffers from the lack of independent observations against which it can be verified.

The major source of uncertainty throughout the year comes from uncertainty in the modelled yield slope  $S$  (see Section 3.4.2); however, in winter, uncertainty from satellite column calculation becomes dominant (see Section 3.4.3). Monthly calculated slope uncertainty mostly lies within 30% to 50%, and for each grid box a month of model data is used, which means that slope uncertainty is not reduced by monthly averaging. Uncertainty from each OMI satellite measurement is relatively large ( $> 100\%$ ); however, averaging thousands of pixels in each grid square greatly reduces the monthly uncertainty. Uncertainty in satellite HCHO is seasonally dependent, with better signal during the summer. Reliable OMI measurements are less frequent (leading to higher uncertainty) at high solar zenith angles, which are more frequent at higher latitudes and during winter. Table 3.10 shows the estimated uncertainty calculated in this work in summer and winter over each region described by Figure 3.17. The assumptions and calculations made to determine uncertainties in the top-down

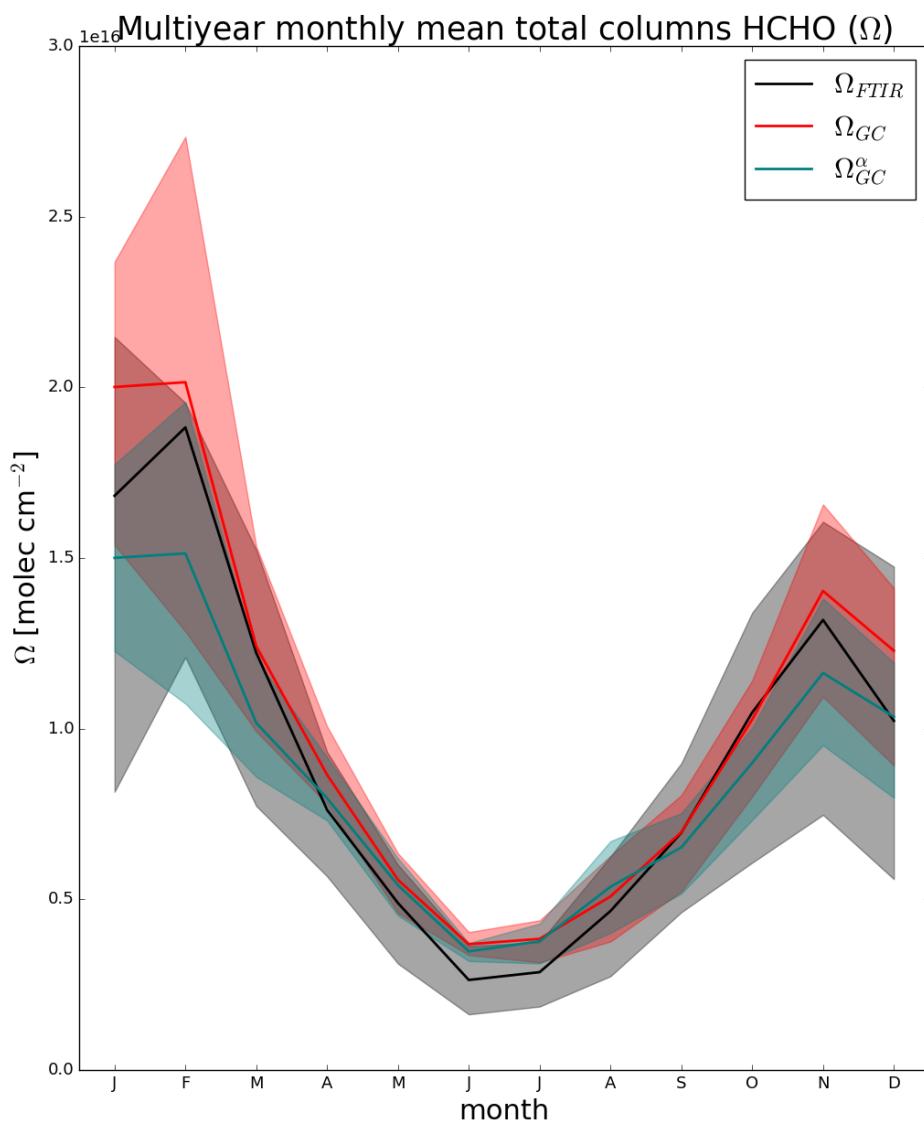


FIGURE 3.35: Multi-year (from all days where both measurements and model data exist: 2007–2012) monthly mean total column ( $\Omega$ ) HCHO from the FTIR instrument at Wollongong (southeast Australia), and the colocated convolved GEOS-Chem equivalent before ( $\Omega_{GC}$ ) and after ( $\Omega_{GC}^\alpha$ ) scaling isoprene emissions. Shaded areas show inter quartile range.

TABLE 3.10: Relative uncertainty estimates.

Region	Summer			Winter		
	$\frac{\Delta E_{OMI}}{E_{OMI}}$	$\frac{\Delta \Omega_{OMI}}{\Omega_{OMI}}$	$\frac{\Delta S}{S}$	$\frac{\Delta E_{OMI}}{E_{OMI}}$	$\frac{\Delta \Omega_{OMI}}{\Omega_{OMI}}$	$\frac{\Delta S}{S}$
Aus	46%	21%	36%	51%	258%	35%
SE	54%	14%	37%	66%	141%	38%
NE	53%	18%	35%	51%	40%	37%
Mid	45%	38%	38%	47%	41%	35%
SW	43%	27%	32%	61%	96%	36%
N	39%	16%	33%	38%	22%	30%

a: Grid squares with monthly uncertainty over 200% are removed when calculating the mean uncertainty, which has small (< 1%) impacts on  $E_{OMI}$  overall, but removes many data points in winter.

estimate ( $\Delta E_{OMI}$ ), the satellite column ( $\Delta \Omega_{OMI}$ ) and the slope ( $\Delta S$ ) are described in the following subsections.

### 3.4.1 Top-down emissions

Important factors in the calculation of isoprene emissions using OMI HCHO include the modelled relationship between HCHO and isoprene and the satellite HCHO measurements. Uncertainty in each of these terms is quantified before being combined in quadrature to give the uncertainty estimate of the a posteriori. Additional biases may arise due to the filters applied to satellite data and model output, and where possible these are assessed.

The final determination of top-down emissions comes from Equation 3.11, repeated here:

$$E_{OMI} = \frac{\Omega_{OMI} - \Omega_{OMI,0}}{S}$$

Assuming each term is independent, the following quadrature rules are used we use the following quadrature rules<sup>mh</sup> to estimate random error in  $E_{OMI}$ :

$$z = x + y : \Delta z = \sqrt{(\Delta x)^2 + (\Delta y)^2} \quad (3.13)$$

$$z = x/y : \Delta z = z \sqrt{\left(\frac{\Delta x}{x}\right)^2 + \left(\frac{\Delta y}{y}\right)^2} \quad (3.14)$$

Which leads to the uncertainty estimation for the a posteriori emissions as follows

$$\Phi \equiv \Omega_{OMI} - \Omega_{OMI,0}$$

$$\Delta\Phi = \sqrt{(\Delta\Omega_{OMI})^2 + (\Delta\Omega_{OMI,0})^2} \quad (3.15)$$

$$\Delta E_{OMI} = E_{OMI} \times \sqrt{\left(\frac{\Delta\Phi}{\Phi}\right)^2 + \left(\frac{\Delta S}{S}\right)^2} \quad (3.16)$$

$\Delta E_{OMI}$  is calculated using the uncertainty in underlying terms:  $\Delta S$ ,  $\Delta \Omega_{OMI}$ , and  $\Delta \Omega_{OMI,0}$ . For  $\Delta S$  ( $\Omega_{GC} = S \times E_{GC} + \Omega_{OMI,0}$  from equation 3.10), the variance in the monthly linear regression of modelled isoprene emissions and column HCHO is used—I use variance in the monthly linear regression of modelled isoprene emissions and column HCHO<sup>mh</sup>, shown in Section 3.4.2. For  $\Delta \Omega_{OMI}$  and  $\Delta \Omega_{OMI,0}$ , uncertainty comes from instrument fitting uncertainty, modelled AMF uncertainty, and uncertainty in the background correction terms, which are described and calculated in Section 3.4.3.

Figure 3.36 shows relative uncertainty over each region of Australia in monthly bins. Uncertainty in the southern regions increases between May and July due to increased error in the satellite measurements. Northern regions are impacted less by the seasonal satellite error, which generally is 10 – 20 % lower than the error from  $S$ . Figure 3.37 shows the spatial distribution of relative uncertainty in the a posteriori in summer and winter. Here the effects of satellite uncertainty at higher latitudes (especially in winter) can be seen through the increase in uncertainty with increasing latitude.

### 3.4.2 Model Uncertainty

$E_{OMI}$  depends partly on the product it is trying to improve, as modelled yield is based on GEOS-Chem run with MEGAN emissions. The uncertainty in the reduced major axis regression slope between model HCHO ( $\Omega_{GC}$ ) and emissions ( $E_{GC}$ ) is used to estimate  $\Delta S$  in Equation 3.16. THere I use the ratio of t<sup>mh</sup>he upper bound of the 95% confidence interval of the regression slope ( $CI_{UB}$ ) is used<sup>mh</sup> to represent the relative uncertainty.

$$\frac{\Delta S}{S} = \frac{CI_{UB}}{S} - 1 \quad (3.17)$$

For example, if the interval upper bound is 30% higher than the slope, relative uncertainty is set to 0.3 (or 30%). The confidence interval for each month is based on the covariance matrix between  $\Omega_{GC}$  and  $E_{GC}$ , and the critical t-statistic considering  $n$  to be the number of days in the month (the significance value of the test is set to 0.025).

This is a simple method of approximating the uncertainty of this term, only accounting for monthly uncertainty of the slope calculation. It does not take into account uncertainty in the underlying model, nor representation uncertainties arising from temporal or spatial resolution, which are difficult to quantify. Figure 3.38 shows the relative uncertainty in  $S$  over Australia and for each region. There is little discernible seasonality to the relative error in  $S$ , which generally ranges from 0.3 – 0.4 (30 – 40%). For comparison, Palmer et al. (2006) found  $\frac{\Delta S}{S}$  to be 30% over the US after comparing with another chemical transport model and in situ measurements. To improve understanding of uncertainty in  $S$  would require further analysis of GEOS-Chem yield over Australia, including how it responds to environmental and meteorological parameters, and how representative this modelled quantity is when compared to measurements. Slope estimation may also be affected by non-isoprene biogenic VOC emissions such as monoterpenes, which are not analysed in this work. The effects of these biogenics will be in part seen by  $\Delta S$  (as they make up some of the variability); however this is an important source of HCHO that will need to be attributed in future work to improve confidence in the estimation of  $S$ .

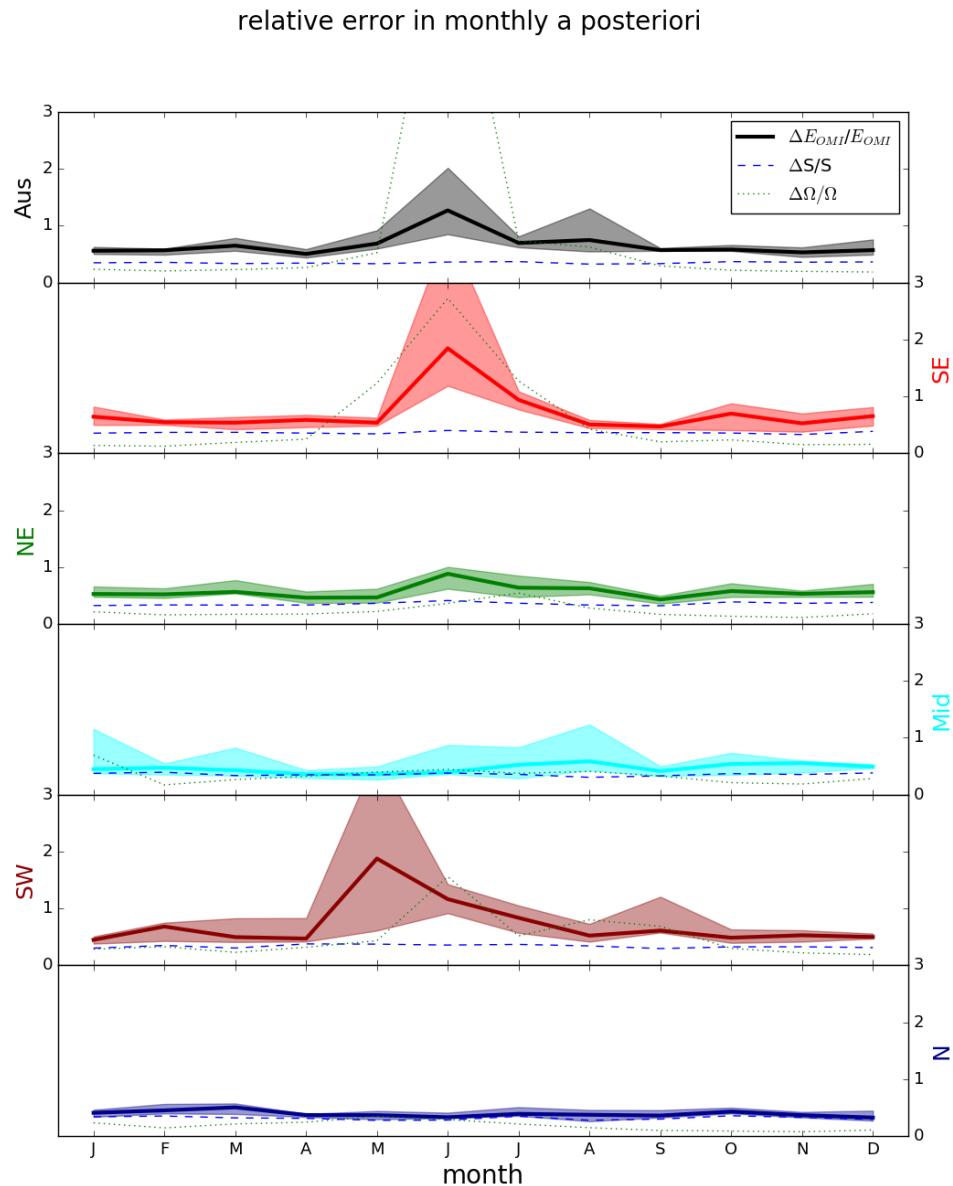


FIGURE 3.36: Median and inter-quartile range of multi-year monthly relative uncertainty in the a posteriori. Median relative uncertainty in  $S$  and  $\Omega$  are added as dashed and dotted lines respectively.

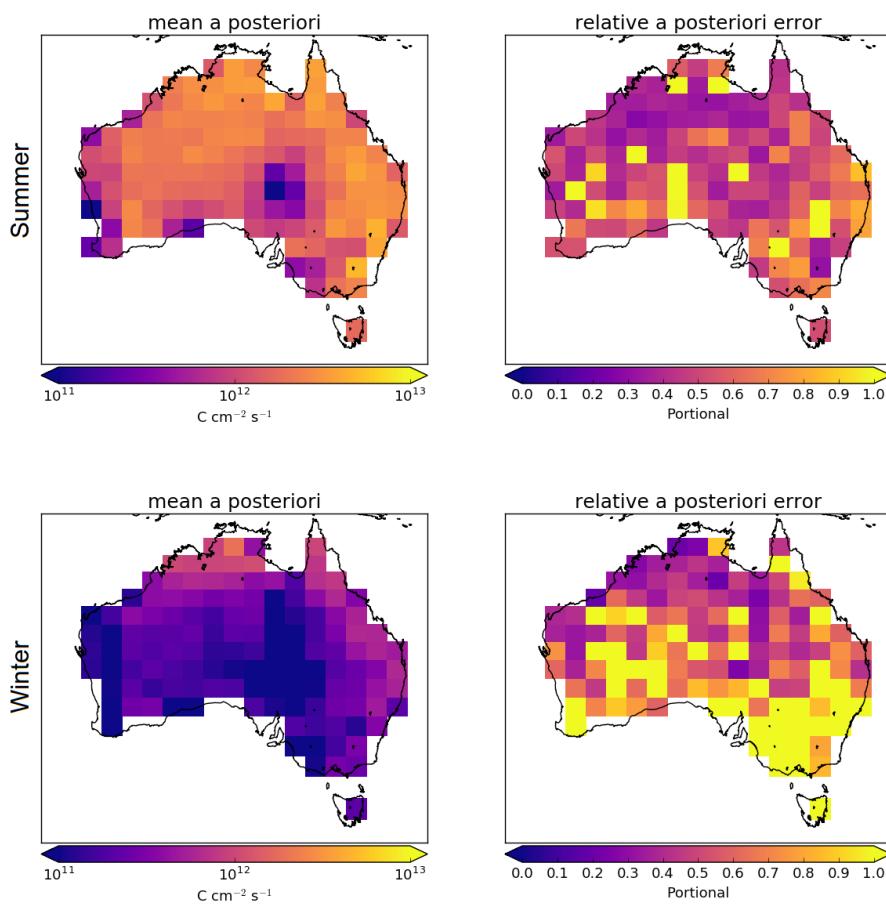


FIGURE 3.37: Summer (DJF, top row) and winter (JJA, bottom row) a posteriori emissions (left column) and relative error (right column).

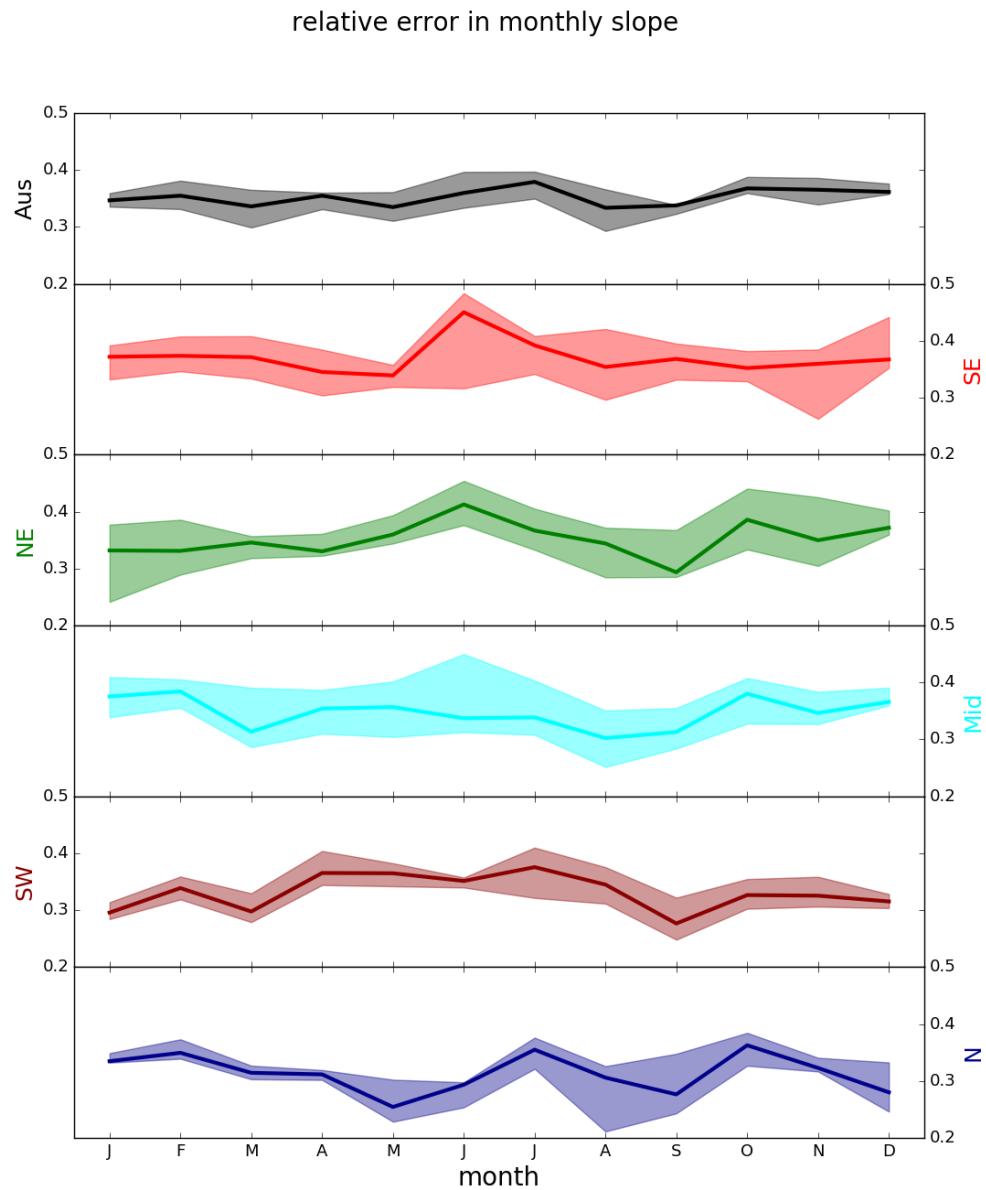


FIGURE 3.38: Median and inter-quartile range of monthly binned uncertainty in  $S$ .

Filtering for spatial smearing (see Section 3.2.6.1) reduces the number of data points making up the regression slope  $S$ . The process generally improves the linear relationship between isoprene emission and HCHO total column, which suggests it is working as intended to remove days when local biogenic emissions are not driving HCHO enhancement. Where  $S$  regressions have a correlation coefficient ( $r$ ) of less than 0.4, a multi-year average (or in the worst cases no value at all) is used in lieu of monthly  $S$  data. The thresholds for the smearing filter are based on literature values from other countries, which may prove to be unsuitable within Australia. To improve the understanding of smearing in Australia, a better approximation of HCHO lifetimes and yields, as well as  $\text{NO}_x$  seasonality and regional concentrations, is required. The filtering process is accounted for in the calculation of  $\Delta S$ , and the effect is a small reduction of the overall uncertainty from  $S$ .

Model biases are not analysed in this thesis, except to note that they would impact both preliminary OMI calculations (via use in calculating the air mass factor) and the modelled slope. Insufficient independent measurements in Australia make it impossible to quantify uncertainty at a national scale.

### 3.4.3 Satellite Uncertainty

Corrected vertical columns of HCHO from the OMI product are calculated using Equation 2.23:  $\Omega = \frac{\text{SC} - \text{RSC}}{\text{AMF}}$ . Error in satellite HCHO columns is determined by error in the three terms  $\text{SC}$ ,  $\text{RSC}$ , and  $\text{AMF}$ :

#### Fitting error from the OMI retrieval

Fitting error represents the uncertainty in the DOAS technique used to estimate HCHO concentrations. Fitting error is provided in the OMHCHO product. This error is ascribed to the  $\text{SC}$  term.

Provided with the OMI product is the measurement of uncertainty in each pixel, calculated by the Smithsonian Astrophysical Observatory from the back scattered solar radiation fit (Gonzalez Abad et al. 2015; Abad et al. 2016). This is used as the  $\Delta \text{SC}$  in Equation 3.18. The relative fitting error per pixel ( $\frac{\Delta \text{SC}}{\text{SC}}$ ) ranges from around 20% to 150%, and is higher where low amounts of HCHO are detected, and at higher solar zenith angles (i.e. at high latitudes).

#### Uncertainty in AMF calculations

Air mass factors model vertical sensitivity to slant path measurements in the satellite instrument, and uncertainties arise predominantly from uncertain cloud parameters (Palmer et al. 2006). Vertical columns from OMI are recalculated using AMFs derived from GEOS-Chem (Section 2.6.3). AMF uncertainty can be determined through comparison of GEOS-Chem output to independently measured HCHO columns. In this work the AMF error is set to 30%, which approximates that seen by Palmer et al. (2006), since measurements over Australia are lacking. For comparison, Wollongong FTIR measurements are over-predicted by GEOS-Chem modelled vertical columns by  $\sim 15 - 20\%$ , but these are not representative at the continental scale simulated here. The assumed error of 30% is ascribed to the  $\text{AMF}$  term.

Palmer et al. (2006) calculated the error in AMF through combining estimates of error in the UV albedo database ( $\sim 8\%$ ), model error based on in situ measurements, cloud error (20 – 30% (Martin 2003)), and aerosol errors ( $< 20\%$ ), totalling AMF error of around  $\sim 30\%$  (calculated in quadrature). This error estimate can be compared with that of Curci et al. (2010), where the error in AMF calculations and background columns were respectively found to be 30% and 15%. Millet et al. (2008) also examined this uncertainty and determine an overall uncertainty ( $1\sigma$ ) of 25 – 27% in HCHO vertical columns with calculated AMFs where cloud fraction  $< 0.2$ .

### Uncertainty of HCHO background

OMI vertical columns are corrected using background (or reference sector) measurements. This is to account for instrument degradation, and adds some uncertainty to the column. In this work vertical columns are corrected using reference sector measurements combined with modelled HCHO, which is described in Chapter 2 (Section 2.6.5). Error from background uncertainty is ascribed to the *RSC* term.

The *RSC*, or background correction, is based on differences in the remote Pacific between daily HCHO slant columns measured by OMI and monthly averages from GEOS-Chem. The correction for each pixel is determined per latitude and OMI track; however, a couple of conservative simplifications are used here to estimate the error in this term. For each SC, the *RSC* is set to the mean correction matching the *SC* latitude over all tracks. For each day, the  $\Delta RSC$  is set to the standard deviation of the *RSC* over Australian latitudes (45°S to 10°S) in all tracks. Background error calculated in this way is on the order of 5%-10% after monthly averaging, however this error increases in the higher southern latitudes during autumn and winter to  $\sim 15\%$ . For comparison, the background error is assumed to be 15% in Curci et al. (2010) following (Dufour et al. 2008).

Calculation of uncertainty in the OMI vertical column HCHO ( $\Omega = \frac{SC - RSC}{AMF}$ ) is performed using quadrature equations 3.13 and 3.14. Error in the slant column ( $\Delta SC$ ) is combined with assumed relative AMF error ( $\frac{\Delta AMF}{AMF}$ ) of 30%, and background error to calculate  $\Delta\Omega_{OMI}$  (and  $\Delta\Omega_{OMI,0}$ ) as follows:

$$\begin{aligned}\Delta(SC - RSC) &= \sqrt{(\Delta SC)^2 + (\Delta RSC)^2} \\ \Delta\Omega &= \Omega \sqrt{\left(\frac{\Delta(SC - RSC)}{(SC - RSC)}\right)^2 + \left(\frac{\Delta AMF}{AMF}\right)^2} \\ \frac{\Delta\Omega}{\Omega} &= \sqrt{\frac{(\Delta SC)^2 + (\Delta RSC)^2}{(SC - RSC)^2} + \left(\frac{\Delta AMF}{AMF}\right)^2}\end{aligned}\quad (3.18)$$

The *RSC* term is described in Chapter 2 Section 2.6.6. Negative columns can occur where column amounts are lower than *RSC*, and these are not removed so as not to introduce a bias. When monthly averages are less than zero, relative error is set to 100%. This only impacts the uncertainty calculations in winter for the non-northern regions, where occasional highly negative absolute uncertainty was seen when  $\Omega$  approached 0.

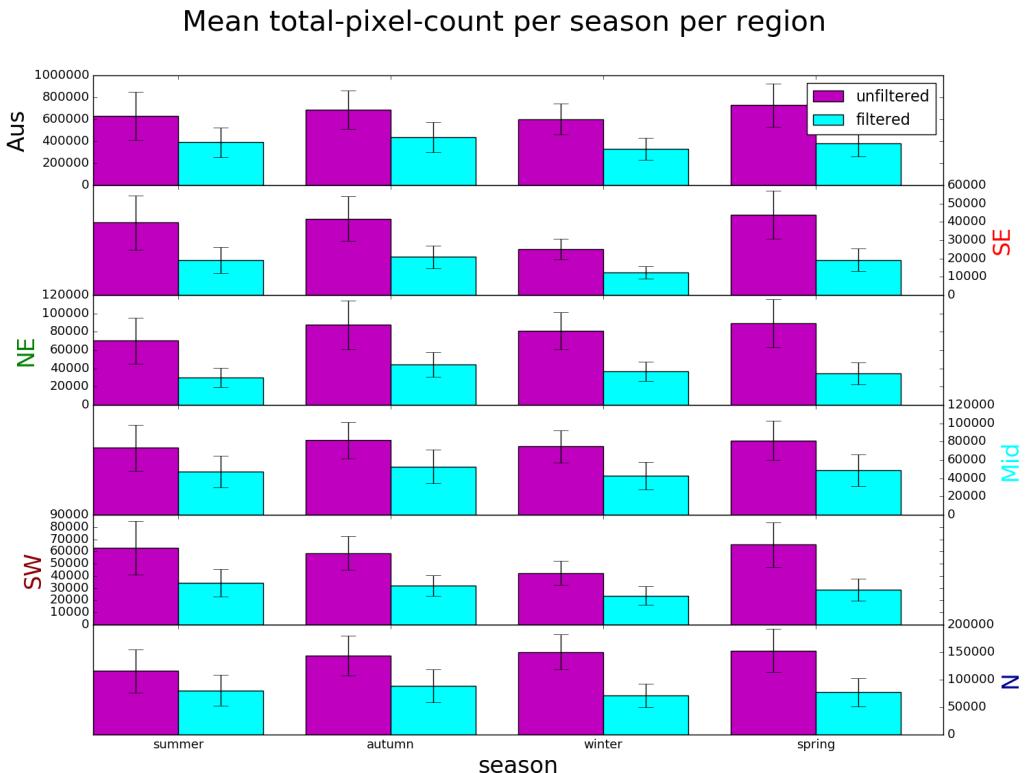


FIGURE 3.39: Mean and standard deviation (vertical error bars) of total pixel counts per region per season, before (magenta) and after (cyan) applying smearing, pyrogenic, and anthropogenic filters.

These sources of error can be reduced through spatial and temporal averaging, as they are assumed to be unbiased. Uncertainty is reduced by the square root of the number of pixels averaged over each  $2^\circ \times 2.5^\circ$  grid square for each day or month. For example, daily averaging reduces pixel uncertainty by a factor of 2-4. Figure 3.39 shows the pixel counts in each region before and after applying filters. Winter has lower pixel count for southern regions, with approximately 50-60% fewer good pixels through May, June, and July. The lowest pixel counts occur in the southeast in winter, likely due to a mix of relatively high solar zenith angles and filtering of anthropogenic emissions. Northern regions have lower pixel counts in the summer, most likely due to increased cloud coverage which limits satellite measurement capabilities as discussed previously.

Figure 3.40 shows the relative uncertainty in monthly satellite columns for each sub-region and averaged over Australia. Uncertainty in winter at higher latitudes is greatly increased due to lower pixel counts, lower absolute column amounts, and higher fitting error.

Uncertainty in satellite HCHO ( $\Delta\Omega$ ) from literature and calculated here is listed in Table 3.11. De Smedt et al. (2012) found satellite HCHO uncertainty to be 30 – 40% for the GOME-2 instrument by combining slant column systematic and random errors. For mid latitude winters they found an excess of 60% uncertainty. OMI measurements

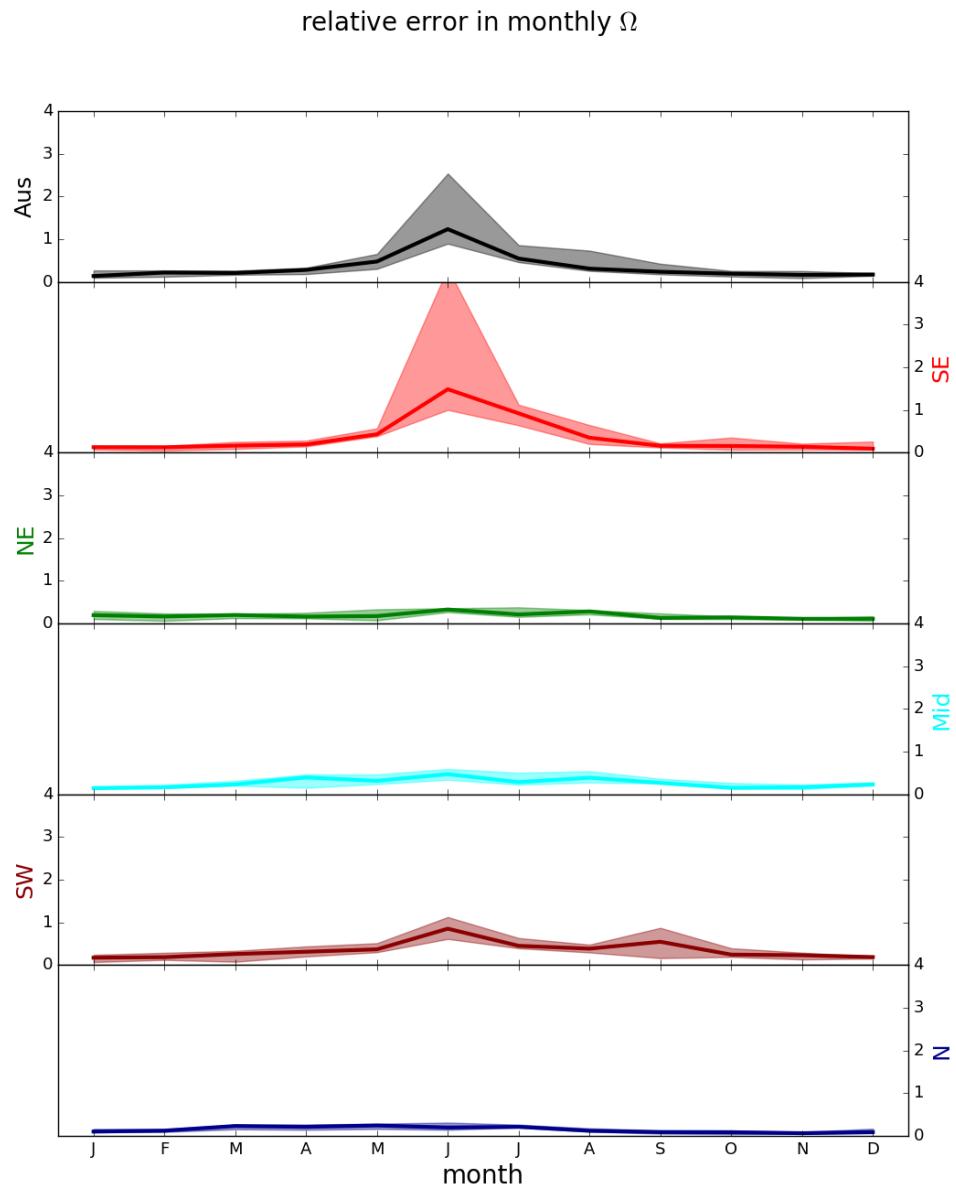


FIGURE 3.40: Median and inter-quartile range for monthly binned relative uncertainty in satellite vertical columns.

TABLE 3.11: Uncertainties in satellite total column HCHO.

uncertainty	location	notes
40%	North America	GOME, mostly due to cloud interference <sup>a</sup>
26%	North America	GOME, OMI, with cloud fraction less than 20% <sup>b</sup>
30%-40%	global	GOME-2 <sup>c</sup>
> 60%	Mid-latitude	GOME-2 in winters <sup>c</sup>
1%-10%	Australia	OMI, monthly uncertainty at $2^\circ \times 2.5^\circ$ <sup>d</sup>
50%-100+%	Australia	OMI, in winter at higher latitudes <sup>d</sup>

a: Millet et al. (2006) and Palmer et al. (2006)

b: Millet et al. (2008)

c: De Smedt et al. (2008) and De Smedt et al. (2012)

d: This work

will have similar uncertainty; however, the array of detectors provide more pixels which can be averaged to reduce this uncertainty.

In order to calculate the bias or systematic error, an understanding of biases in the underlying terms is required, since there is little in the way of comparable measurements. OMI has been shown to underestimate observed HCHO by up to  $\sim 40\%$  elsewhere in the world. For example OMI underestimated aircraft measurements by 37% in Guyana (Barkley et al. 2013). OMI underestimates range from 20-37% when compared against aircraft data over the southeast United States (Zhu et al. 2016). OMI validation against 7 sites (the most southern site being Reunion Island at  $20.9^\circ\text{S}$ ) using MAX-DOAS and FTIR retrievals showed up to 50% underestimates by satellite HCHO retrievals (De Smedt et al. 2015). The highest underestimate occurred during periods of high concentration (Vigouroux et al. 2009; De Smedt et al. 2015). Satellite HCHO may also suffer from  $\sim 13\%$  overestimation when taking monthly averages due to only measuring on relatively cloud-free days (Sur, Palmer, and Abad 2018). In this thesis, the a posteriori is linearly related to the satellite HCHO and any bias is directly transferred. The conclusion drawn here is that the isoprene emissions product in this work may be affected by satellite HCHO underestimation of up to 40%, and also by monthly HCHO overestimation of 13%, which gives a potential bias of 1/0.6 to 1/1.13. This may be complicated further if the satellite bias over Australia does not match the bias over the remote Pacific at corresponding latitudes. However, bias over Australia cannot be quantified due to insufficient measurements. GEOS-Chem biases would affect the recalculation of HCHO, but they cannot be quantified and so are not included in this thesis. Lacking suitable measurements to estimate satellite bias over Australia, the potential bias range compiled from the literature ( $-13\%$  to  $+40\%$ ) is applied to the mean emissions in each month when calculating the uncertainty range shown in the results (see Figure 3.20).

### 3.4.4 Sensitivity to AMF recalculation

This section examines the sensitivity of the top-down isoprene emission estimates ( $E_{OMI}$ ) to the AMF recalculation method. The a posteriori emissions change linearly with recalculated vertical columns, which are calculated in three different ways: using

the AMF provided in the OMHCHO product ( $AMF_{OMI}$ ), recalculating AMF shape factors but keeping the original scattering weights ( $AMF_{GC}$ ), or recalculating both shape factors and scattering weights ( $AMF_{PP}$ ), as described in Section 2.5.

Figure 3.41 shows AMFs and emissions recalculated using each of these three methods over 2005. This figure only depicts grid squares with non-zero emissions, since in this work the regions of interest are those where emissions are relatively more substantial within Australia. The fully recalculated  $AMF_{PP}$  is higher in summer, but lower in other seasons compared against the other recalculations. The direct effect of a lower AMF is an increased vertical column, which should lead to higher emission estimates, and vice versa. This can be seen in all seasons except for summer, although even here the emissions estimates based on  $AMF_{PP}$  are higher than if they are calculated using the original  $AMF_{OMI}$ . Emission estimates vary widely over Australia, but the sensitivity to AMF recalculation technique is proportional and potentially non-linear. Changes in AMF of 5 – 30% cause changes in emission estimates of 5 – 50%. Further analysis is warranted and should be a focus of future work.

### 3.4.5 Sensitivity to filtering

Figure 3.42 shows emissions estimates with and without filtering for smearing, anthropogenic, and pyrogenic influences. The overall effect of filtering is to slightly raise emissions in all non-summer months, with relatively little change to the mean in winter months. This is true in all regions except for northern Australia, which shows a slight decrease in spring. Eastern regions are most frequently filtered, with more than 50% of the available good pixels filtered throughout the year, while other regions vary between 20% and 60% with lower filtering rates in summer months.

## 3.5 Conclusions and implications

Very few ground-based measurements of BVOC concentrations are available in Australia. Emission models use largely unverified extrapolations for emission factors and parameterisations in Australia, leading to overestimated isoprene emissions (Emmerman et al. 2016). This leads to uncertainty and error when modelling atmospheric ozone and other trace gases.

In this chapter, I created and tested<sup>mh</sup> an isoprene emissions estimate (a posteriori) is created and tested<sup>mh</sup> based on OMI satellite measurements of HCHO, a high-yield product of isoprene oxidation. The a posteriori emissions' effects on ozone and HCHO were tested by running GEOS-Chem with a seasonal (multi-year monthly averaged) gridded ( $2^\circ \times 2.5^\circ$ ) scaling factor calculated from the a posteriori applied to the a priori emissions. Uncertainty in the primary components of the top-down emissions calculation were calculated where possible and potential biases identified.

The a posteriori isoprene emission estimate showed that GEOS-Chem coupled with the MEGAN emissions model overestimates emissions in summer by a factor of 2-5. Total yearly Australian emissions are reduced from  $39 \text{ Tg yr}^{-1}$  to  $21 \text{ Tg yr}^{-1}$  (decrease of  $\sim 46\%$ ) in the a posteriori. The overestimate is spatially and temporally diverse and leads to model biases in HCHO, and ozone. Running GEOS-Chem using

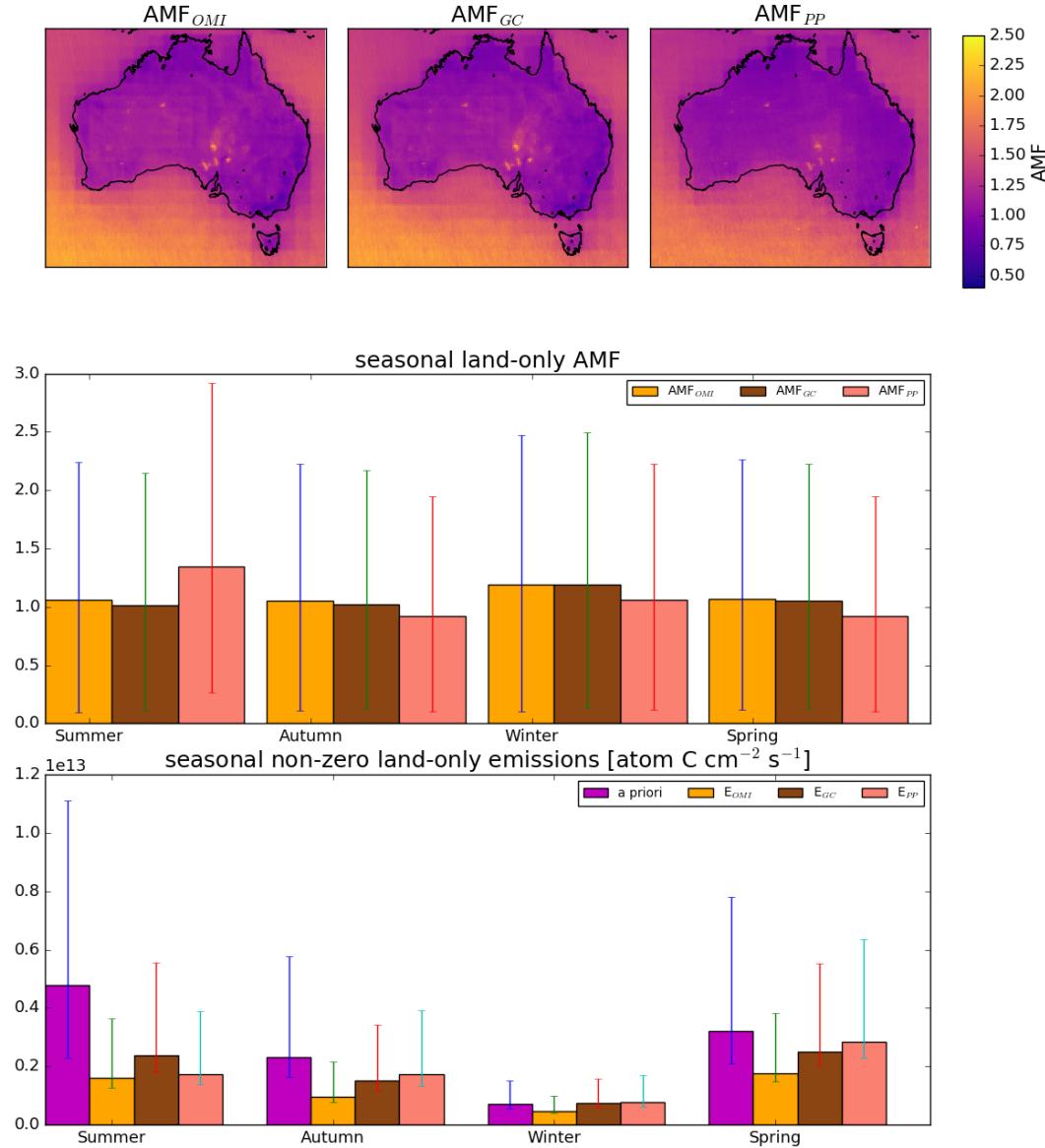


FIGURE 3.41: Top row: averaged OMI Satellite AMF for 2005, from the OMHCCHO dataset (left,  $AMF_{OMI}$ ), recalculated using GEOS-Chem shape factors (middle,  $AMF_{GC}$ ), and recalculated using GEOS-Chem shape factors and scattering weights (right,  $AMF_{PP}$ ). Middle row: mean and inter-quartile range of each AMF over 2005 for each season. Bottom row: mean and inter-quartile range of non-zero a priori emissions based on the three AMFs (with matching subscripts) along with the a priori emissions from GEOS-Chem.

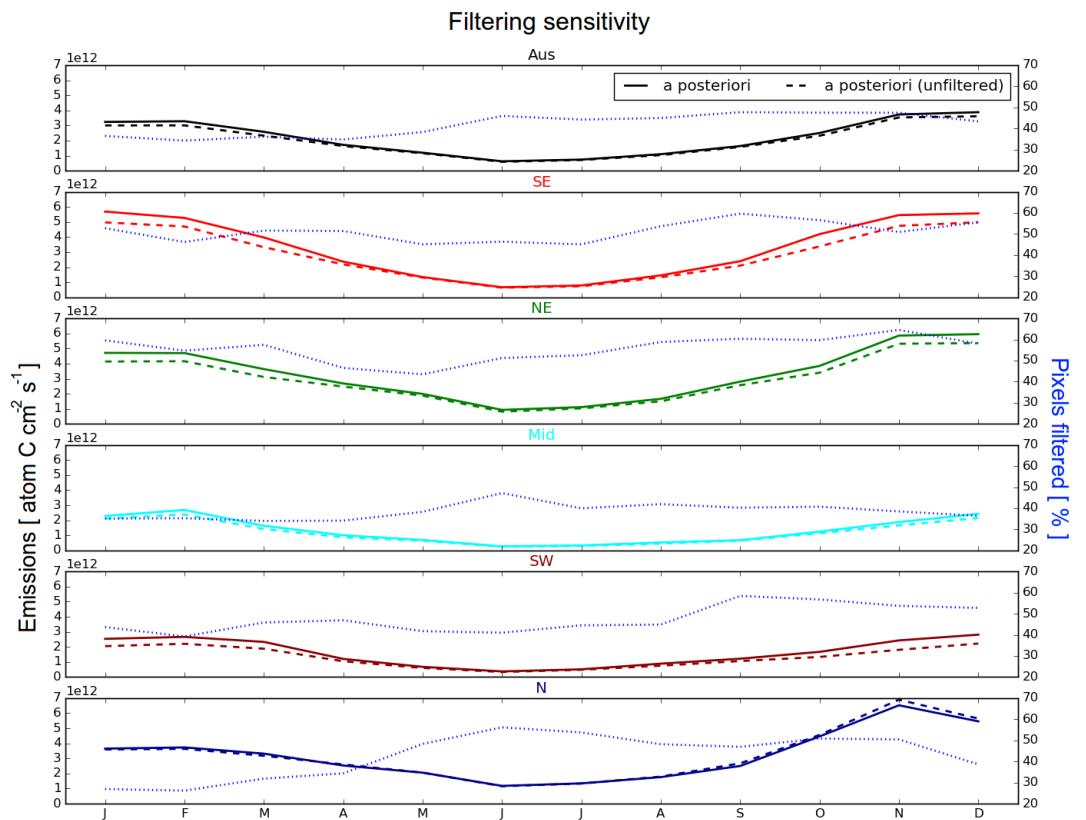


FIGURE 3.42: Multi-year monthly mean values for a posteriori emission estimates calculated with (solid) and without (dashed) applying filters for anthropogenic, pyrogenic, and smearing influences. The portion of pixels within each region which are filtered is shown on the right axis with a blue dotted line.

scaled emissions based on OMI HCHO columns reduced the model HCHO overestimate (when compared to OMI HCHO) in summer by half, from  $\sim 100\%$  down to  $\sim 50\%$ , with most of the difference occurring outside the northern region. Model vertical column HCHO variance is somewhat ( $\sim 10\% - 50\%$ ) lower than that seen by the OMI satellite, and this difference increased after scaling isoprene emissions. Scaling GEOS-Chem emissions also lowered simulated surface ozone concentrations by  $\sim 5\%$ . A posteriori uncertainty was shown to be on the order of 50% (monthly, per grid square), with large satellite-based uncertainty in winter and a potential bias coming from satellite data of  $-13\%$  to  $+40\%$ . The primary uncertainty in the a posteriori emissions comes from the monthly modelled isoprene-to-HCHO yield ( $\sim 30\% - 50\%$  uncertainty), although at higher latitudes in winter the satellite uncertainty becomes restrictively high ( $> 100\%$ ).

The reason for the a priori overestimate remains unclear, and both global and Australian emissions estimates for isoprene range widely. The bias is in part due to the MEGAN emission model's parameterisation of how isoprene responds to parameters such as leaf area index, plant specific emission factors, and meteorological factors. In Australia, a mixture of poorly defined emission factors (e.g., Emmerson et al. 2016), unaccounted for soil moisture forcing (e.g., Sindelarova et al. 2014; Emmerson et al. 2019) and poorly understood forest responses to meteorological stresses likely drive uncertainty and model biases (Jiang et al. 2018; Emmerson et al. 2019). A potential improvement to the MEGAN soil parameterisation has been described (Jiang et al. 2018); however, a detailed map of soil properties is not currently available in Australia, and this would be required to apply the parameterisation here. An alternative parameterisation of soil moisture and drought effects on isoprene emissions was recently implemented in a different CTM by Emmerson et al. (2019) and applying this parameterisation in GEOS-Chem should be a priority for future work in Australia.

In the US, bias between OMI and in situ measurements is as high as 40%, but bias across Australia cannot be determined as there are not enough independent observations. This leads to a wide uncertainty range for estimated top-down emissions (see Figure 3.20) and limits potential top-down refinements to isoprene emissions. Ground-based and aircraft VOC,  $\text{NO}_x$ , HCHO, and ozone measurements over large areas at relatively fine temporal resolution would help quantify the currently unknown satellite biases while additionally providing constraints for bottom-up models. In the northern region in particular, emissions are affected by monsoonal forcing, but increased cloud coverage during the monsoon limits satellite coverage. This makes characterisation of forest emissions and their response to sunlight, temperature, and moisture even more important in these areas. In addition to measurements, further analysis determining the sensitivity of modelled emissions to model resolution and changing soil moisture parameters (and parameterisations) would provide the foundation to improve GEOS-Chem and MEGAN modelled isoprene emissions along with oxidation products like HCHO and ozone.

In the future, other satellites (e.g., GOME-2, TROPOMI) could be used to improve emission estimates further, with differing overpass times potentially allowing a measure of diurnal emission patterns. Additionally, an adjoint version of GEOS-Chem over Australia could provide improved estimates of isoprene emission taking into account transport. An adjoint based inversion could also provide an evaluation of

how resolution-limited the linear top-down emission estimates are over Australia. The emission estimate created in this chapter could also be refined to higher temporal resolution, with further analysis of uncertainty. Oversampling techniques could be applied near populated areas in order to improve the understanding of isoprene, HCHO, and ozone relationships over cities (e.g., Surl, Palmer, and Abad 2018). Furthermore, linking the top-down isoprene emissions to the underlying emission factors for Australia and implementing the changes in GEOS-Chem would improve the understanding of the natural atmosphere over this relatively remote portion of the planet.

### 3.6 Contributions and acknowledgements

Model runs, data retrieval, analysis and error calculations and attribution in this Chapter are performed by the Author, with feedback and direction provided by Jenny Fisher and Clare Paton-Walsh. AMF and satellite corrections are performed by the Author, following discussions with Chris Miller, Paul Palmer, Luke Surl, and Gonzalez Abad.<sup>tb</sup>



## Chapter 4

# Stratospheric ozone intrusions

### 4.1 Foreword

This chapter is unchanged (excepting titles, labels, and numbering) from a paper on which I was the first author, published in the journal of Atmospheric Chemistry and Physics: Jesse W Greenslade et al. (2017). “Stratospheric ozone intrusion events and their impacts on tropospheric ozone in the Southern Hemisphere”. In: *Atmospheric Chemistry and Physics*, pp. 1–33. URL: <https://www.atmos-chem-phys.net/17/10269/2017/acp-17-10269-2017.pdf>. Additionally this chapter mentions a supplementary document, which has been added to this thesis as Appendix C. Contributions (Section 4.9) are additionally duplicated here along with an estimate of what proportion of the work that was performed by the author.<sup>mh</sup>

Jesse Greenslade wrote the algorithms, ran the GEOS-Chem simulations, performed the analysis and led the writing of the paper under the supervision and guidance of Simon Alexander, Robyn Schofield, and Jenny Fisher. Andrew Klekociuk contributed the Davis ozonesonde data and performed the analysis of the alternate STT proxy. All authors contributed to editing and revising the manuscript. While more than 95% of the work was performed by the Author, none of it would be possible without the assistance, advice, and expertise of each of the coauthors and those mentioned in the acknowledgements (Section 4.9).<sup>mh</sup>

### 4.2 Introduction

Tropospheric ozone constitutes only 10% of the total ozone column but is an important oxidant and greenhouse gas which is toxic to life, harming natural ecosystems and reducing agricultural productivity. Over the industrial period, increasing tropospheric ozone has been estimated to exert a radiative forcing (RF) of  $365 \text{ mWm}^{-2}$  (Stevenson et al. 2013), equivalent to a quarter of the CO<sub>2</sub> forcing (Forster et al. 2007). While much tropospheric ozone is produced photochemically from anthropogenic and natural precursors, downward transport from the ozone-rich stratosphere provides an additional natural source of ozone that is particularly important in the upper troposphere (Jacobson and Hansson 2000, and references therein). The contribution of this source to overall tropospheric ozone budgets remains uncertain (Škerlav, Sprenger, and Wernli 2014), especially in the southern hemisphere (SH). Models show that stratospheric ozone depletion has propagated to the upper troposphere (Stevenson et al. 2013). However, work based on the Southern Hemisphere Additional

OZonesonde (SHADOZ) network suggests stratospheric mixing may be increasing upper tropospheric ozone near southern Africa (Liu et al. 2015; Thompson et al. 2014). Uncertainties in the various processes which produce tropospheric ozone limit predictions of future ozone-induced radiative forcing. Here we use a multi-year record of ozonesonde observations from sites in the southern hemisphere extra-tropics, combined with a global model, to better characterise the impact of stratospheric ozone on the tropospheric ozone budget in the southern hemisphere.

Stratosphere-to-troposphere transport (STT) primarily impacts the ozone budget in the upper troposphere but can also increase regional surface ozone levels above the legal thresholds set by air quality standards (Danielsen 1968; Lelieveld et al. 2009; Lefohn et al. 2011; Langford et al. 2012; Zhang et al. 2014; Lin et al. 2015). In the western US, for example, deep STT events during spring can add 20-40 ppbv of ozone to the ground-level ozone concentration, which can provide over half the ozone needed to exceed the standard set by the U.S. Environmental Protection Agency (Lin et al. 2012; Lin et al. 2015). Another hotspot for STT is the Middle East, where surface ozone exceeds values of 80 ppbv in summer, with a stratospheric contribution of 10 ppb (Lelieveld et al. 2009). Estimates of the overall contribution of STT to tropospheric ozone vary widely (e.g. Galani 2003; Stohl et al. 2003; Stevenson et al. 2006; Lefohn et al. 2011). Early work based on two photochemical models showed that 25-50% of the tropospheric ozone column can be attributed to STT events globally, with most contribution in the upper troposphere (Stohl et al. 2003). In contrast, a more recent analysis of the Atmospheric Chemistry and Climate Model Inter-comparison Project (ACCMIP) simulations by Young et al. (2013) found that STT is responsible for  $540 \pm 140 \text{ Tg yr}^{-1}$ , equivalent to  $\sim 11\%$  of the tropospheric ozone column, with the remainder produced photochemically (Monks et al. 2015). This wide range in model estimates exists in part because STT is challenging to accurately represent, and finer model resolution is necessary to simulate small scale turbulence. Observation-based process studies are therefore key in determining the relative frequency of STT events, with models then able to quantify STT impact over large regions. Ozonesondes are particularly valuable for this purpose as they provide multi-year datasets with high vertical resolution.

Lower stratospheric and upper tropospheric ozone concentrations are highly correlated, suggesting mixing across the tropopause mainly associated with the jet streams over the Atlantic and Pacific oceans (Terao et al. 2008). Extra-tropical STT events most commonly occur during synoptic-scale tropopause folds (Sprenger, Croci Maspoli, and Wernli 2003; Tang and Prather 2012; Frey et al. 2015) and are characterised by tongues of high potential vorticity (PV) air descending to lower altitudes. As these tongues become elongated, filaments disperse away from the tongue and mix irreversibly into the troposphere. STT can also be induced by deep overshooting convection (Frey et al. 2015), tropical cyclones (Das et al. 2016) and mid-latitude synoptic scale disturbances (e.g. Stohl et al. 2003; Mihalikova et al. 2012). STT events have been observed in tropopause folds around both the polar front jet (Vaughan, Price, and Howells 1993; Beekmann et al. 1997) and the subtropical jet (Baray et al. 2000). The summertime pool of high tropospheric ozone over the eastern Mediterranean (EM) is mainly attributed to the downward ozone transport, as a result of the enhanced subsidence (Zanis et al. 2014) and the tropopause fold activity (Akritidis et al. 2016) over the

region. The EM exhibits a summer maximum of subsidence, which sits between 20°E and 35°E and 31°N to 39°N, while zonally most subtropical tropopause folds occur during winter (Tyrlis et al. 2014, and references therein). They are also observed near cut-off lows (Price and Vaughan 1993; Wirth 1995), so both regional weather patterns and stratospheric mixing are important to understand for STT analysis.

Stratospheric ozone intrusions undergo transport and mixing, with up to half of the ozone diffusing within 12 hours following descent from the upper troposphere (Trickl et al. 2014). The long range transport of enhanced ozone can be facilitated by upper tropospheric winds, with remarkably little convective mixing, as shown by Trickl et al. (2014) who measure STT air masses two days and thousands of kilometres from their source. Cooper et al. (2004) shows how STT advection can transport stratospheric air over long distances, with a modelled STT event spreading from the northern Pacific to the East coast of the USA over a few days.

The strength (ozone enhancement above background levels), horizontal scale, vertical depth, and longevity of these intruding ozone tongues vary with wind direction and strength, topography, and season. While the frequency, seasonality, and impacts of STT events have been well described in the tropics and northern hemisphere (NH), observational estimates from the SH extra-tropics are noticeably absent in the literature. The role of STT in the SH remains highly uncertain due to the more limited data availability compared to the NH and the temporal sparsity of these datasets (Mze et al. 2010; Thompson et al. 2014; Liu et al. 2015).

Here, we characterise the seasonal cycle of STT events and quantify their contribution to the SH extra-tropical tropospheric ozone budget using nearly a decade of ozonesonde observations from three locations around the Southern Ocean spanning latitudes from 38°S-69°S. In Section 4.3 we describe the observations and methods used to identify STT events and to relate STT occurrence to meteorological events. We show how possible biomass burning smoke plume influence is detected and handled, and we introduce the GEOS-Chem model which is used for ozone flux estimation. Within Section 4.4 we show the seasonality, altitude, depth, and frequency of detected STT events, along with a comparison of our findings to other literature where possible. In Section 4.5 we analyse how well GEOS-Chem captures the tropospheric ozone seasonality and quantity near our three sites. In Section 4.6 an extrapolation of the STT detection frequencies along with GEOS-Chem monthly tropospheric ozone columns is used to estimate STT ozone flux near our three sites. We also compare and contrast our results against relevant literature. Finally, in Section 4.7 we examine in detail the uncertainties involved in our STT event detection technique and ozone flux estimations.

## 4.3 Data and Methods

### 4.3.1 Ozonesonde record in the Southern Ocean

Ozonesondes provide a high vertical resolution profile of ozone, temperature, pressure, and humidity from the surface and up to 35 km. In the troposphere, the ozonesondes generally perform 150-300 measurements. Ozone mixing ratio is quantified with

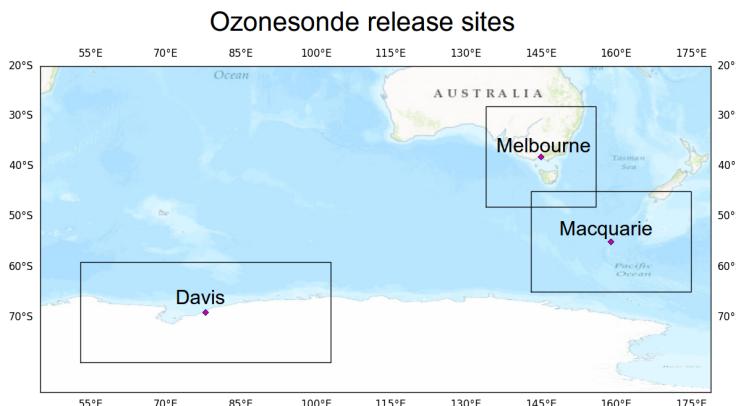


FIGURE 4.1: Ozonesonde release sites and the regions used to examine STT effect on tropospheric ozone levels.

TABLE 4.1: Number of sonde releases at each site over the period of analysis.

Site	Total Releases	Monthly Releases (J, F, M, ...)	Date Range
Davis	240	11, 12, 13, 12, 17, 31, 29, 28, 32, 28, 15, 12	2006/04/13 - 2013/11/13
Macquarie Island	390	32, 31, 45, 28, 34, 33, 28, 35, 29, 33, 31, 31	2004/01/20 - 2013/01/09
Melbourne	456	31, 38, 40, 38, 41, 36, 38, 39, 46, 40, 38, 31	2004/01/08 - 2013/12/18

an electrochemical concentration cell, using standardised procedures when constructing, transporting, and releasing the ozonesondes (<http://www.ndsc.ncep.noaa.gov/organize/protocols/appendix5/>). Ozonesondes are estimated to provide around 2% precision in the stratosphere, which improves at lower altitudes, and ozonesondes have been shown to be accurate to within 5% when the correct procedures are followed (Smit et al. 2007).

Ozonesondes are launched approximately weekly from Melbourne (38° S, 145° E), Macquarie Island (55° S, 159° E) and Davis (69° S, 78° E), as shown in Fig. 4.1. Melbourne is a major city in the south east of Australia, and may be affected by anthropogenic pollution in the lower troposphere. Macquarie Island is isolated from the Australian mainland, situated in the remote Southern Ocean and unlikely to be affected by any local pollution events. Davis is on the coast of Antarctica and also unlikely to experience the effects of anthropogenic pollution.

For this study, we use the 2004-2013 data for Melbourne and Macquarie Island and the 2006-2013 data for Davis because both ozone and geopotential height (GPH) are available from the World Ozone and Ultraviolet Data Centre archived data in these periods. At Davis, ozonesondes are launched twice as frequently during the ozone hole season and preceding months (June-October) as at other times of year (Alexander et al. 2013). A summary of ozonesonde releases at each site can be seen in Table 4.1.

Characterisation of STT events requires a clear definition of the tropopause. Two common tropopause height definitions are the standard lapse rate tropopause (WMO

1957) and the ozone tropopause (Bethan, Vaughan, and Reid 1996). The lapse rate tropopause is defined as the lowest altitude where the lapse rate (vertical gradient of temperature) is less than  $2^{\circ}\text{C km}^{-1}$ , provided the lapse rate averaged between this altitude and 2 km above is also below  $2^{\circ}\text{C km}^{-1}$ . The ozone tropopause is defined as the lowest altitude satisfying the following three conditions for the ozone mixing ratio (OMR) (Bethan, Vaughan, and Reid 1996):

1. Vertical gradient of OMR is greater than  $60 \text{ ppb km}^{-1}$ ;
2. OMR is greater than 80 ppb; and
3. OMR exceeds 110 ppb between 500 m and 2000 m above the altitude under inspection (modified to between 500 m and 1500 m in the Antarctic, including the site at Davis).

The ozone tropopause may misdiagnose the real tropopause altitude during stratosphere-troposphere exchange; however, it is useful at polar latitudes in winter, where the lapse-rate definition may result in artificially high values for tropopause height (Bethan, Vaughan, and Reid 1996; Tomikawa, Nishimura, and Yamanouchi 2009; Alexander et al. 2013). We require lapse rate defined tropopauses to be at a minimum of 4 km altitude. Another commonly used tropopause definition is determined with the use of PV (dynamical tropopause). In the extra-tropics the isosurface where  $\text{PV} = 2 \text{ PVU}$  ( $1 \text{ PVU} = 10^{-6} \text{ m}^2 \text{ s}^{-1} \text{ K kg}^{-1}$ ) is often used to define the tropopause, allowing the 3D representation of tropopause folds and other tropopause features in a sufficiently resolved model (Škerlak, Sprenger, and Wernli 2014; Tyrlis et al. 2014). The PV is not calculable using the ozonesonde measurements alone, so in this work the ozone tropopause is used when determining STT events or measured tropopause altitude.

Figure 4.2 shows the monthly median ozone tropopause altitudes at each location (solid lines). At Melbourne, the tropopause altitude displays a seasonal cycle with maximum in summer and minimum in winter. This seasonality is missing at Macquarie Island and almost reversed at Davis, which has a minimum during autumn and maximum from winter to spring. Tropopause altitude decreases with latitude from 9-14 km at Melbourne ( $38^{\circ}$  S) to 7-9 km at Davis ( $69^{\circ}$  S).

Figure 4.3 shows multi-year averaged ozone mixing ratios measured by ozonesonde over the three stations. Over Melbourne, increased ozone extending down through the troposphere is apparent from December to March and from September to November. The increased tropospheric ozone in these months is due to STT (in summer), and possible biomass burning influence (in spring), both discussed in more detail in the following sections. Over Davis and Macquarie Island, tropospheric ozone is higher between March and October, although the seasonal differences are small compared to those at Melbourne. The seasonality shown in Fig. 4.3 for Davis is consistent with remote free tropospheric photochemistry determined by solar radiation availability and temperature, resulting in higher ozone in winter (Lelieveld and Dentener 2000). NO<sub>2</sub> stratospheric observations have been conducted in the Southern hemisphere at Lauder, Macquarie Island and Arrival Heights (i.e. Struthers et al. 2004) which displays a winter minima in seasonality consistent with an ozone maxima. Influence from the ozone hole can be seen over Davis in October, with relatively low ozone levels extending up 5-6 km into the stratosphere.

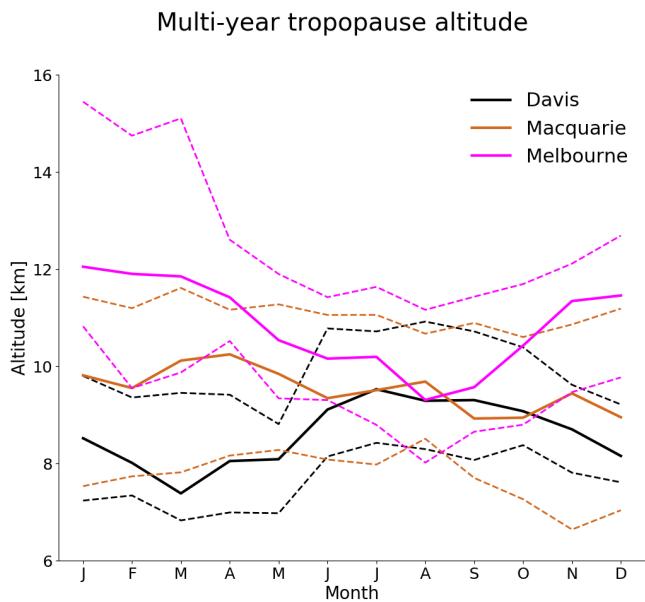


FIGURE 4.2: Multi-year monthly median tropopause altitude (using the ozone defined tropopause) determined from ozonesondes measurements at Davis (2006-2013), Macquarie Island (2004-2013), and Melbourne (2004-2013) (solid lines). Dashed lines show the 10th to the 90th percentile of tropopause altitude for each site.

### 4.3.2 Model description

To provide regional and global context to the ozonesonde observations, we use the GEOS-Chem version 10-01 global chemical transport model (Bey et al. 2001), which simulates ozone along with more than 100 other trace gases throughout the troposphere and stratosphere. Stratosphere-troposphere coupling is calculated using the stratospheric unified chemistry extension (UCX) (Eastham, Weisenstein, and Barrett 2014). Transport is driven by assimilated meteorological fields from the Goddard Earth Observing System (GEOS-5) maintained by the Global Modeling and Assimilation Office (GMAO) at NASA. Ozone precursor emissions are from the Model of Emissions of Gases and Aerosols from Nature (MEGAN) version 2.1 (Guenther et al. 2012) for biogenic emissions, the Emissions Database for Global Atmospheric Research (EDGAR) version 4.2 for anthropogenic emissions, and the Global Fire Emissions Database (GFED4) inventory (Giglio, Randerson, and Van Der Werf 2013) for biomass burning emissions. Our simulation was modified from the standard v10-01 to fix an error in the treatment of ozone data from the Total Ozone Mapping Spectrometer (TOMS) satellite used to calculate photolysis (see [http://wiki.seas.harvard.edu/geos-chem/index.php/FAST-JX\\_v7.0\\_photolysis\\_mechanism#Fix\\_for\\_TOMS\\_to\\_address\\_strange\\_cycle\\_in\\_OH\\_output.](http://wiki.seas.harvard.edu/geos-chem/index.php/FAST-JX_v7.0_photolysis_mechanism#Fix_for_TOMS_to_address_strange_cycle_in_OH_output.)).

Our simulations span 2005-2012 (following a 1-year spin-up) with horizontal resolution of  $2^{\circ}$  latitude by  $2.5^{\circ}$  longitude and 72 vertical levels from the surface to 0.01 hPa. The vertical resolution is finer near the surface at  $\sim 60$  m between levels,

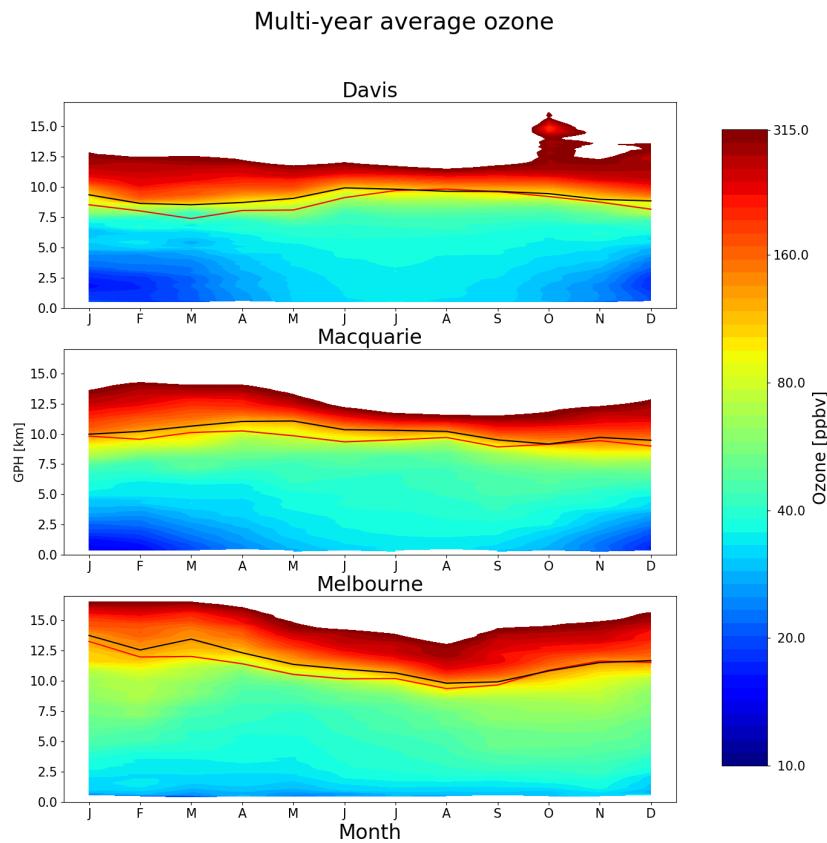


FIGURE 4.3: Multi-year mean seasonal cycle of ozone mixing ratio over Davis, Macquarie Island, and Melbourne as measured by ozonesondes. Measurements were interpolated to every 100 m and then binned monthly. Black and red solid lines show median ozone and lapse-rate defined tropopause altitudes (respectively), as defined in the text.

spreading out to  $\sim 500$  m near 10 km altitude. For comparison to the ozonesonde observations, the model state was saved every 6 hours within the grid boxes containing each site. When comparing against ozonesondes, GEOS-Chem UTC+0 time samples are used for all sites. This means that the simulated ozone profiles are analysed at local times of 7AM for Davis, and 11AM for Macquarie Island and Melbourne. GEOS-Chem uses the tropopause height provided by GEOS-5 meteorological fields, which are calculated using a lapse-rate tropopause definition using the first minimum above the surface in the function  $0.03 \times T(p) - \log(p)$ , with  $p$  in hPa (Rienecker 2007).

#### 4.3.3 Characterisation of STT events and associated fluxes

We characterise STT events using the ozonesonde vertical profiles to identify tropospheric ozone enhancements above a local background (in moles per billion moles of dry air, referred to here as ppb). The process is illustrated in Figure 4.4 on an example ozone profile. First, the ozone vertical profiles are linearly interpolated to a regular

grid with 20 m resolution from the surface to 14 km altitude. Small vertical-scale fluctuations in ozone, which are captured by the high-resolution ozonesondes, can be regarded as sinusoidal waves superimposed on the large vertical scale background tropospheric ozone. As such, the interpolated profiles are bandpass-filtered using a fast Fourier transform (Press et al. 1992) to retain these small vertical scales, between 0.5 km and 5 km (removing low and high frequency perturbations). The high frequency perturbations are removed as they may represent noise in the measurements. The perturbations with scales longer than 5 km represent the vertical gradient of ozone concentration from the surface to the stratosphere. In what follows, these filtered vertical profiles are referred to as perturbation profiles.

For an event to qualify as STT, a clear increase above the background ozone level is needed, as a bandpass filter leaves us with enhancements minus any noise or seasonal scale vertical profile effects. We next use all the perturbation profiles at each site to calculate the 95th percentile perturbation value for the site. The threshold is calculated from all the interpolated filtered values between 2 km above the surface and 1 km below the tropopause. This is our threshold for tropospheric ozone perturbations, and any profiles with perturbations exceeding this value in individual ozonesondes are classified as STT events. STT events at altitudes below 4 km are removed to avoid surface pollution, and events within 0.5 km of the tropopause are removed to avoid false positives induced by the sharp transition to stratospheric air. We note that this ozone detection methodology detailed above does not allow us to resolve STT events where the ozone flux is spread diffusely across the troposphere without a peak-like structure in the ozonesonde profile. In other words, STT events which might have occurred some distance and time away from the location of the ozonesonde profiles may not be readily detected using the high vertical resolution, but infrequent, ozonesonde launches.

We define the ozone peak as the altitude where the perturbation profile is greatest between 2 km from the surface and 0.5 km below the tropopause. The STT event is confirmed if the perturbation profile drops below zero between the ozone peak and the tropopause, as this represents a return to non-enhanced ozone concentrations. Alternatively, the STT event is also confirmed if the OMR between the ozone peak and the tropopause drops below 80 ppb and is at least 20 ppb lower than the OMR at the ozone peak. If neither of these conditions are met, the profile is rejected as a non-event. This final step removes near-tropopause anomalies for which there is insufficient evidence of detachment from the stratosphere. Vertical ozone profiles recorded by ozonesondes are highly dependent on the time of launch (Sprenger, Croci Maspoli, and Wernli 2003), and it cannot be guaranteed that detected ozone enhancements are fully separated from the stratosphere, although this method minimises that risk by removing detected events too near the tropopause.

We estimate the ozone flux into the troposphere associated with each event by integrating the ozone concentration enhancement vertically over the altitude range for which an STT event is identified (i.e. enhancement near the ozone peak over which the perturbation profile is greater than zero). This estimate is conservative because it does not take into account any ozone enhancements outside of the detected peak that may have been caused by the STT, and also ignores any enhanced ozone background amounts from synoptic-scale stratospheric mixing into the troposphere.

### Ozone at Melbourne on 2004/01/08

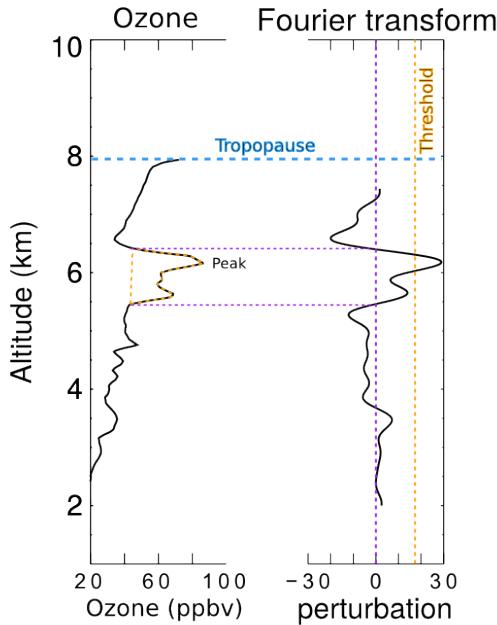


FIGURE 4.4: An example of the STT identification and flux estimation methods used in this work. The left panel shows an ozone profile from Melbourne on 8 January 2004 from 2 km to the tropopause (blue dashed horizontal line). The right panel shows the perturbation profile created from bandpass filtering of the mixing ratio profile. The STT occurrence threshold calculated from the 95th percentile of all perturbation profiles is shown as the orange dashed line, and the vertical extent of the event is shown with the purple dashed lines (see details in text). The ozone flux associated with the STT event is calculated using the area outlined with the orange dashed line in the left panel.

Our method differs somewhat from that used by Tang and Prather (2010) to detect STT events from ozonesonde measurements. Their definition is based on subjective analysis of sondes released from 20 stations ranging in latitude from 35° S to 40° N. They identify an STT event if, starting from 5 km altitude, ozone exceeds 80 ppb and then within 3 km decreases by 20 ppb or more to a value less than 120 ppb. Their technique would miss many events due to the lower ozone concentrations found in the cleaner Southern Hemisphere.

#### 4.3.4 Biomass burning influence

The STT detection algorithm described in Sect. 4.3.3 assumes all ozone enhancements are caused by stratospheric intrusions. In some cases, however, these perturbations may in fact reflect ozone production in lofted smoke plumes. Biomass burning in southern Africa and South America has previously been shown to have a major influence on atmospheric composition in the vicinity of our measurement sites (Oltmans et al. 2001; Gloudemans et al. 2006; Edwards et al. 2006), particularly from July to

December (Pak et al. 2003; Liu et al. 2017a). On occasion, smoke plumes from Australian and Indonesian fires can also reach the mid-high southern latitudes, as seen from satellite measurements of carbon monoxide (CO) discussed below.

Large biomass burning events emit substantial quantities of ozone precursors, some of which are capable of being transported over long distances and driving ozone production far from the fire source (Jaffe and Wigder 2012). Ozone production from biomass burning is complex and affected by photochemistry, fuel nitrogen load, and time since emission, among other factors. While ozone production occurs in some biomass burning plumes, this is not always the case; therefore ozone perturbations detected during transported smoke events may or may not be caused by the plume. For this reason all detected STT events which could be caused by smoke plumes are flagged, following the procedure outlined below. Calculations of seasonality, and ozone flux do not include flagged events, however they are included in summary plots in this work.

Possible biomass burning influence is identified using satellite observations of CO from the AIRS (Atmospheric Infra-red Sounder) instrument on board the Aqua satellite (Texeira 2013). CO is emitted during incomplete combustion and is an effective tracer of long-range transport due to its long lifetime (Edwards 2003; Edwards et al. 2006). In the Southern Hemisphere, biomass burning is the primary source of CO, making CO a good proxy for fire plumes (e.g. Sinha et al. 2004; Mari et al. 2008). To identify possible biomass burning influence, AIRS vertical column CO is visually inspected for all dates with detected STT events. Smoke plumes are diagnosed over areas with elevated CO columns ( $\sim 2 \times 10^{18}$  molecules  $\text{cm}^{-2}$  or higher), and any sonde-detected STT event that occurs near (within  $\sim 150$  km of) a smoke plume is flagged. Removal of these detections reduces the yearly estimated ozone flux by  $\sim 15\%$  at Macquarie Island and  $\sim 20\%$  at Melbourne.

All days with detected STT events were screened, with the exception of one event during which there were no available AIRS data (January 2010). We find that biomass burning may have influenced 27 events over Melbourne and 21 events over Macquarie Island. These events are flagged in the following sections, and are not used in our calculation of total STT flux. All of the flagged events except for two occurred during the SH burning season (July to December). No events at Davis were seen to be influenced by smoke transport.

### 4.3.5 Classifying synoptic conditions during STT events

Synoptic scale weather patterns are examined using data from the European Centre for Medium-range Weather Forecasts (ECMWF) Interim Reanalysis (ERA-I) (Dee et al. 2011). This is done using the ERA-I data products over the three sites on dates matching the detected STT events. We use the ERA-I 500 hPa data to subjectively classify the events based on their likely meteorological cause, by visually examining each date where an event was detected. During STT occurrence, the upper troposphere is typically characterised by nearby cyclones, cut-off lows, or cold fronts. Over Melbourne and Macquarie Island, we find that frontal and low pressure activity are prevalent during STT events (see Sect. 4.4). Over Davis, the weather systems are often less clear, however we see a higher portion of probable cut-off lows. The stratospheric polar

TABLE 4.2: Total number of ozonesonde detected STT events, along with the number of events in each category (see text).

Site	Events	Cut-offs	Frontals	Misc	Fire
Davis	80	44	19	17	0
Macquarie Island	105	19	31	34	21
Melbourne	127	28	31	41	27

vortex may create tropopause folds without other sources of upper tropospheric turbulence such as low pressure fronts or cyclones (e.g. Baray et al. 2000; Sprenger, Croci Maspoli, and Wernli 2003; Tyrlis et al. 2014). Cut-off low pressure systems can be seen clearly in synoptic scale weather maps as regions with lowered pressure and cyclonic winds. Low pressure fronts in the higher southern latitudes travel from west to east and lower the tropopause height. We examine two cases in detail to illustrate the relationship between synoptic-scale conditions and STT events over Melbourne. These are included in a supplementary document (Fig. S2 and S3) which show an archetypal cut-off low and low pressure front. To detect cut-off low pressure systems we look for cyclonic winds and a detached area of low pressure within  $\sim 500$  km of a site on days of event detection. For low pressure fronts we look for low pressure troughs within  $\sim 500$  km. Frontal passage is a known cause of STT as stratospheric air descends and streamers of ozone-rich air break off and mix into the troposphere (Sprenger, Croci Maspoli, and Wernli 2003).

#### 4.4 STT event climatologies

Figure 4.5 shows the seasonal cycles of STT frequency at Davis, Macquarie Island, and Melbourne. Frequency is determined as detected event count divided by total launched ozonesondes for each month. STT events in Figures 4.5-4.8 are coloured based on the meteorological classification described in Sect. 4.3.5, with events classified as either low pressure fronts (“frontal”, dark blue), cut-off low pressure systems (“cutoff” teal), or indeterminate (“misc”, cyan). Events that may have been influenced by transported smoke plumes (Sect. 4.3.4) are shown in red. Ozonesonde releases are summarised in Table 4.1 and detected event counts are summarised in Table 4.2.

There is an annual cycle in the frequency of STT events (Fig. 4.5) with a summertime peak at all three sites. This summertime peak is due to a prevalence of summer low-pressure storms and fronts, which increase turbulence and lower the tropopause (Reutter et al. 2015). At Davis, there are more STT detections during winter relative to our other sites, which may be due to the polar vortex and its associated lowered tropopause and increased turbulence. STT events associated with cut-off low pressure systems are more prevalent during summer, while STT events associated with frontal passage occur throughout the year. The high frequency of STT ozone enhancements is comparable to the  $> 25\%$  frequencies seen over Turkey and east of the Caspian sea in an ERA-I analysis performed by Tyrlis et al. (2014).

The SH summer maximum we see for STT ozone flux can also be seen in Fig. 16 of Škerlak, Sprenger, and Wernli 2014, which shows seasonal flux over the southern ocean, although this is less clear over Melbourne. This seasonality is not clear in the

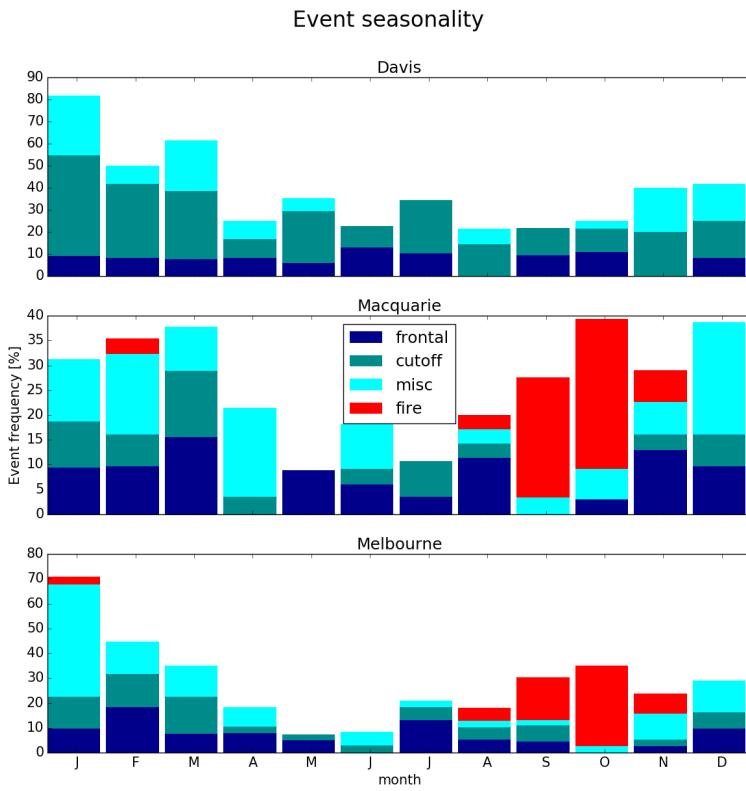


FIGURE 4.5: Seasonal cycle of STT event frequency at Davis (top), Macquarie Island (middle), and Melbourne (bottom). Events are categorised by associated meteorological conditions as described in the text, with low pressure fronts (“frontal”) in dark blue, cut-off low pressure systems (“cutoff”) in teal, and indeterminate meteorology (“misc”) in cyan. Events that may have been influenced by transported smoke plumes are shown in red (see text for details).

recent ERA-Interim tropopause fold analysis performed by Škerlak et al. (2015), where a winter maximum of tropopause fold frequency ( $\sim 0.5\%$  more folds in winter) over Australia can be seen to the north of Melbourne. Their work seems to show slightly higher fold frequencies over Melbourne in summer (Škerlak et al. 2015, Fig. 5), however not to the same extent that our summer peak suggests. Their winter maximum is in the subtropics only - from around  $20^\circ$  S to  $40^\circ$  S, which can be seen as the prevalent feature over Australia in their Fig. 5. Wauben, Fortuin, and Velthoven 1998 look at modelled (CTM driven by ECMWF output) and measured ozone distributions and find more SH ozone in the lower troposphere during austral winter, however they note that the ECMWF fields are uncertain here again due to lack of measurements. Their work shows a generally cleaner lower troposphere in the SH summer but this can not be construed to suggest more or less STT folds in either season. Sprenger, Croci Maspoli, and Wernli 2003 examine modelled STT folds using ECMWF output over March 2000 - April 2001, and show that for this year there is a clear austral winter maximum, again over the  $20^\circ$  S to  $40^\circ$  S band. The winter maximum does not

include Melbourne, or the southern ocean, which explains why we see a seasonality not readily evident in these global-scale studies.

The measurement sites are not in the regions which have a clear winter maximum seen in Fig. 1 Sprenger, Croci Maspoli, and Wernli 2003, and the large scale winter maximum shown by all three studies seems to be dominated by the system in that region. The seasonality of our three sites is not driven by the larger STT system seen over the southern Indian ocean and middle Australia which dominates prior analysis near or over Australia.

To examine the robustness of the distributions shown in Fig. 4.5, we developed an alternative assessment of the seasonal occurrence of STT events, with results shown in Fig. 4.6. Here STT occurrence is evaluated by consideration of the square of the dry Brunt-Väisälä frequency ( $N^2$ ) at the heights of the ozone tropopause ( $z_{OT}$ ) and lapse rate tropopause ( $z_{LRT}$ ) in each ozonesonde profile that has been binned to 100 m resolution. We use  $N^2$  to assess atmospheric stability, which is normally distinctly higher in the stratosphere than in the troposphere, and assume that the vertical temperature gradients within the intrusion respond most rapidly to transported heat, which is an additional characteristic of stratospheric air.  $N^2$  is evaluated using 250 m resolution data (to smooth variability in the vertical gradient of potential temperature that is due to small temperature fluctuations likely associated with gravity waves). The altitude binning chosen is a compromise between vertical resolution and the level of variability in  $N^2$  introduced by temperature gradients associated with perturbations from gravity waves and changes near the lapse rate tropopause, and is the minimum that produces a robust seasonal distribution. We define STT as having taken place if  $N^2(z_{OT}) > N^2(z_{LRT})$  and  $z_{OT} < z_{LRT}$ ; in this way the characteristically higher static stability and ozone concentration of stratospheric intrusion is regarded as being retained as it penetrates below the lapse rate tropopause. The seasonal distributions shown for the three stations in Fig. 4.6 are generally similar to those shown in Fig. 4.5 (although detected events are less frequent), with the main exception that very few events are identified with the alternative method at Davis in the first half of the year. For our STT proxy, we only detect intrusions where the lowest altitude of the intrusion satisfies the ozone tropopause definition. During summer and autumn, the vertical ozone gradients at Davis are weaker compared with the other seasons, and the detected ozone tropopause tends to lie above the lapse rate tropopause potentially reducing the ability to identify STT events based on the definition of our proxy.

Figure 4.7 shows the altitudes of detected events, based on the altitude of peak tropospheric ozone (local maximum ozone within enhancement altitude) in the ozonesonde profile. STT event peaks most commonly occur at 6–11 km above Melbourne and anywhere from 4–9 km at Davis and Macquarie Island. There is no clear relationship between meteorological conditions and event altitude, which may reflect the fact that the ozonesondes observe a snapshot of an event at different stages of its life cycle.

Figure 4.8 shows the distance from the event peak to the ozone defined tropopause, referred to as event depth. The majority of STT events occur within 2.5 km of the tropopause at Davis and Macquarie Island. Over Melbourne, the event depth is more spread out, with peak ozone enhancement generally occurring up to 6 km below the tropopause. Again, there is no clear relationships between meteorological conditions and event depth.

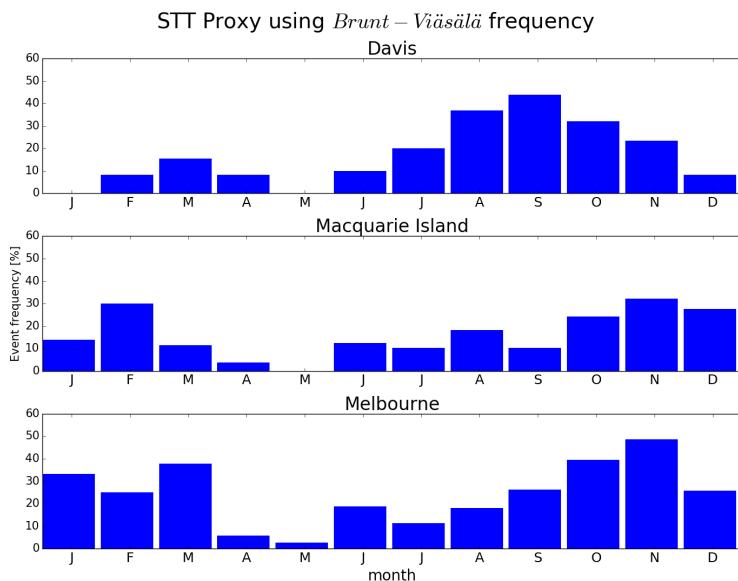


FIGURE 4.6: Seasonal distribution of STT events using the alternative STT proxy, obtained from consideration of the static stability at the ozone and lapse rate tropopauses, for Davis (2006-2013), Macquarie Island (2004-2013), and Melbourne (2004-2013).

## 4.5 Simulated ozone columns

Figure 4.9 compares the time series of tropospheric ozone columns ( $\Omega_{O_3}$ ) in molecules  $\text{cm}^{-2}$  simulated by GEOS-Chem (red) to the measured tropospheric ozone columns (black). GEOS-Chem outputs ozone density ( $\text{molecules cm}^{-3}$ ), and height of each simulated box, as well as which level contains the tropopause, allowing modelled  $\Omega_{O_3}$  to be calculated as the product of density and height summed up to the box below the tropopause level. In both observations and model, the maximum ozone column at Melbourne occurs in austral summer, with a minimum in winter, while Macquarie Island and Davis show the opposite seasonality.

GEOS-Chem provides a reasonable simulation of the observed seasonality and magnitude of  $\Omega_{O_3}$ . Reduced major axis regression of observed versus simulated  $\Omega_{O_3}$  gives a line of best fit with slopes of 1.08 for Davis, 0.99 for Macquarie Island, and 1.34 for Melbourne. The model is only partially able to reproduce the variability in the observations, with paired  $r^2$  values of 0.38 for Davis, 0.18 for Macquarie Island, and 0.37 for Melbourne. Much of the variability is driven by the seasonal cycle, and after removing this effect (by subtracting the multi-year monthly means), the  $r^2$  values decrease to 0.07, 0.11, and 0.30 respectively, although the slope improves at Melbourne to 1.08.

Figure 4.10 shows the observed and simulated ozone profiles at all sites, averaged seasonally. The model generally underestimates ozone in the lower troposphere (up to 6 km) over Davis, although this bias is less pronounced during summer. Over Melbourne, ozone in the lower troposphere is well represented, but the model overestimates ozone from around 4 km to the tropopause. Over Macquarie Island we see

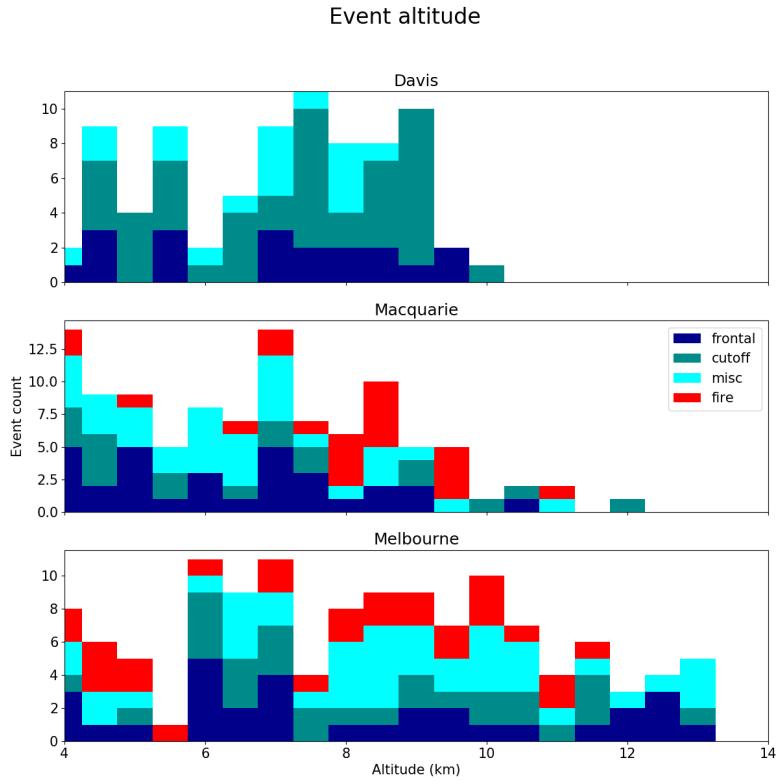


FIGURE 4.7: The distribution of STT events' altitudes at Davis (top), Macquarie Island (middle), and Melbourne (bottom), determined as described in the text. Events are coloured as described in Fig. 4.5.

model overestimation of ozone above 4 km, as well as underestimated ozone in the lower troposphere, suggesting that this region is influenced by processes seen at both of our other sites. Also shown is the mean tropopause height simulated by the model (horizontal dashed red line), which is always higher than the observed average, although this difference is not statistically significant. The effect of local pollution over Melbourne during austral summer (DJF) can be seen from the increased mean mixing ratios and enhanced variance near the surface. The gradient of the O<sub>3</sub> profiles is steeper in the measurements than the model, at all sites during all seasons. Recently Hu et al. 2017 examined GEOS-Chem ozone simulations and found a similar overestimation of upper troposphere ozone in the mid southern latitudes when using the GEOS5 meteorological fields.

Figure 4.11 compares modeled (red) and observed (black) ozone profiles on three example days when STT events were detected using the ozonesondes. The figures show the profile for each site with the closest (qualitative) match between model and observations. The resolution (both vertical and horizontal) of GEOS-Chem in the upper troposphere is too low to consistently allow detection of STTs, although in a few cases (e.g., Melbourne in Fig. 4.11) it appears that the event was large enough to be visible in the model output.

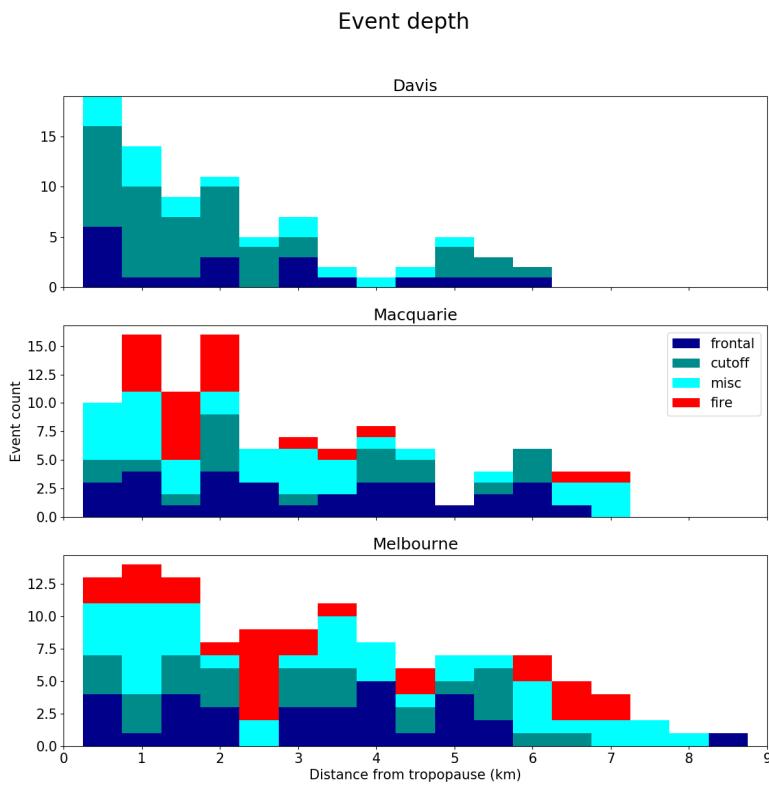


FIGURE 4.8: The distribution of STT events' depths, defined as the distance from the event to the tropopause, at Davis (top), Macquarie Island (middle), and Melbourne (bottom), determined as described in the text.

Events are coloured as described in Fig. 4.5.

## 4.6 Stratosphere-to-troposphere ozone flux from STT events

### 4.6.1 Method

We quantify the mean stratosphere-to-troposphere ozone flux due to STT events at each site based on the integrated ozone amount associated with each STT event (see Sect. 4.3.3). Events that may have been influenced by transported biomass burning are excluded from this calculation. Our estimate provides a preliminary estimate of how much ozone is transported from the stratosphere by the events detected by our method. The estimate is conservative for several reasons: it ignores secondary ozone peaks which may also be transported from the stratosphere, it ignores potential ozone enhancements which may have dispersed and increased the local background mixing ratio, and any influence from STT events nearby which may also increase the local background ozone.

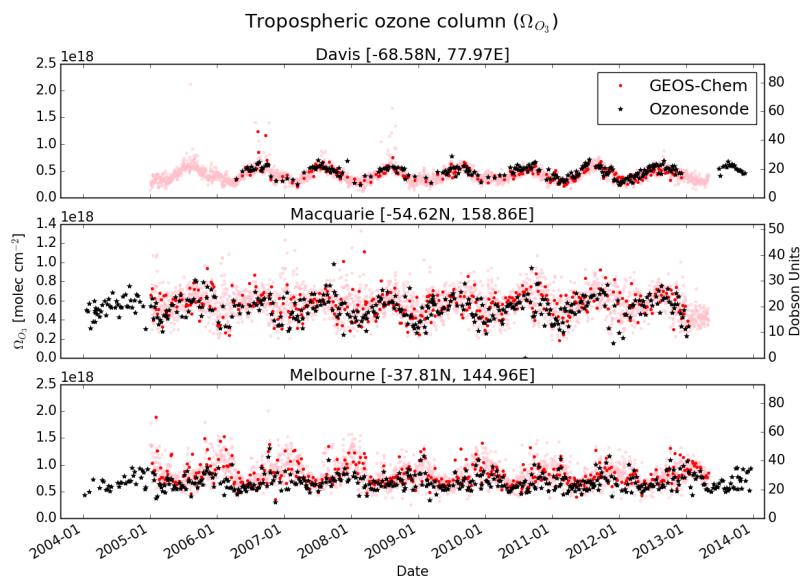


FIGURE 4.9: Comparison between observed (black) and simulated (pink, red) tropospheric ozone columns ( $\Omega_{O_3}$ , in molecules  $\text{cm}^{-2}$ ) from 1 January 2004 to 30 April 2013. For the model, daily output is shown in pink, while output from days with ozonesonde measurements are shown in red. For each site, the model has been sampled in the relevant grid square.

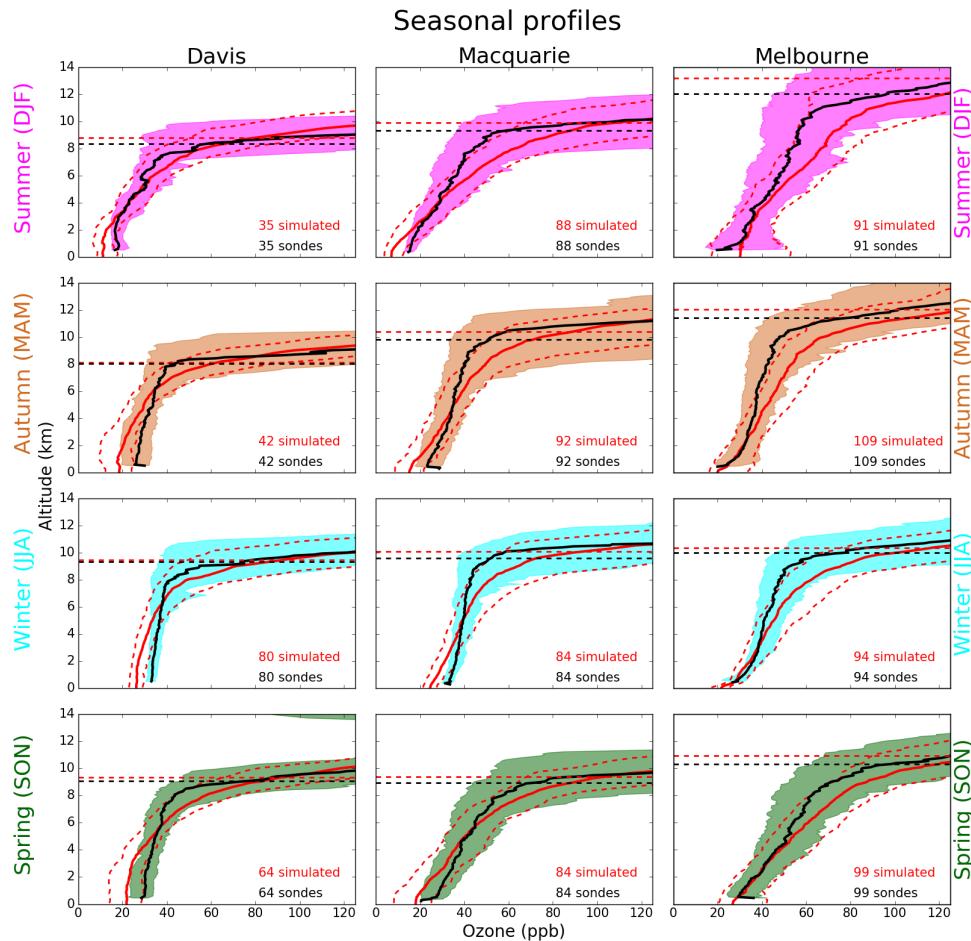


FIGURE 4.10: Observed and simulated tropospheric ozone profiles over Davis, Macquarie Island, and Melbourne, averaged seasonally. Model medians (2005–2013 average) are shown as red solid lines, with red dashed lines showing the 10th and 90th percentiles. Ozonesonde medians (over each season, for all years) are shown as black solid lines, with coloured shaded areas showing the 10th and 90th percentiles. The horizontal dashed lines show the median tropopause heights from the model (red) and the observations (black).

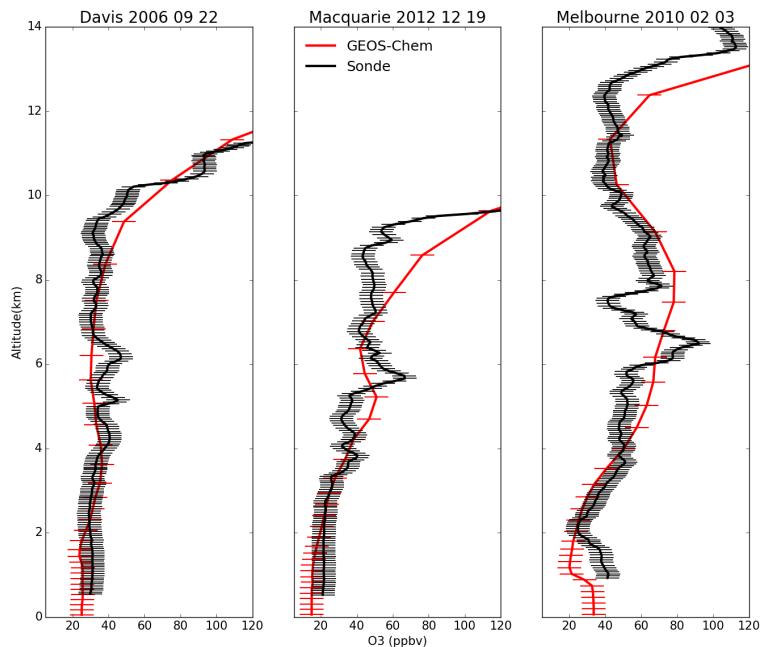


FIGURE 4.11: Example comparisons of ozone profiles from ozonesondes (black) and GEOS-Chem (red) from three different dates during which STT events were detected from the measurements. The dates were picked based on subjective visual analysis. The examples show the best match between model and observations for each site. GEOS-Chem and ozonesonde pressure levels are marked with red and black dashes respectively.

Observed tropospheric columns are calculated from the ozonesondes by calculating the ozone number density (molecules cm<sup>-3</sup>) using measured ozone partial pressure ( $P_{O_3}$ ) and integrating vertically up to the tropopause:

$$\Omega_{O_3} = \int_0^{TP} \frac{P_{O_3}(z)}{k_B \times T(z)} dz$$

where  $z$  is the altitude (GPH),  $TP$  is the altitude at the tropopause,  $T$  is the temperature, and  $k_B$  is the Boltzmann constant.

Three regions are used to examine possible STT flux over a larger area using modeled tropospheric ozone concentrations. The regions are shown in Fig. 4.1. The regions are centred at each site, plus or minus ten degrees latitude, and plus or minus 25, 16, and 11 degrees longitude for Davis, Macquarie Island, and Melbourne respectively. These boundaries approximate a rectangle centred at each site with  $\sim 2000$  km side lengths, covering  $\sim 4.4$ ,  $4.6$ , and  $4.8$  million square km, for Davis, Macquarie Island, and Melbourne respectively.

To determine the ozone column attributable to STT, we determine monthly averaged STT impact ( $I$ ; fraction of tropospheric ozone sourced from the stratosphere as shown above) and the monthly mean tropospheric ozone column (from the GEOS-Chem multi-year mean,  $\Omega_{O_3}$ ) over the regions described above. This can be expressed simply as the STT flux per event (flux<sub>*i*</sub> in each month:  $\text{flux}_i = \Omega_{O_3} \times I$ ). Next we determine how many events are occurring per month by assuming only one event can occur at one time, and that no event is measured twice. These assumptions allow a simple estimate of events per month from the relatively sparse dataset and should hold true as long as our regions of extrapolation are not too large. The ( $P$ )robability of any sonde launch detecting an event is calculated as the fraction of ozonesonde releases for which an STT event was detected, calculated for each month. We assume events last  $N$  days, then find how many events per month we expect by multiplying the days in a month by  $P$  and dividing by this assumed event lifetime. For example if we expect to see an event 25% of the time in a month, and events last one day, we expect one event every four days ( $\sim 7.5$  events in that month) whereas if we expect events to last a week then we would expect  $\sim$ one event in that month. This leads us to multiply our  $\text{flux}_i$  by  $P$ , and then by the term  $M$  ( $M = \frac{\text{days per month}}{N}$ ) determined by our assumed event lifetime in order to determine monthly STT ozone flux.

The longevity of ozone events is very difficult to determine, and we have chosen 2 days as a representative number based on several examples in Lin et al. 2012 where intrusions were seen to last from 1-3 days (occasionally longer) and an analysis of one large event by Cooper et al. 2004 showing that most of the ozone had dispersed after 48 hours. Worth noting is the recent work of Trickl et al. 2014, where intrusions are detected  $> 2$  days and thousands of kilometres away from their initial descent into the troposphere over Greenland or the Arctic. In those regions with high wind shear, mixing appears to be slower, which allows ozone intrusions to be transported further without dissipating into the troposphere. Relative uncertainty in our  $M$  term is set to 50%, as we assume these synoptic events to generally last from 1-3 days.

TABLE 4.3: Seasonal STT ozone contribution in the regions near each site, in  $\text{kg km}^{-2} \text{ month}^{-1}$ . In parentheses are the relative uncertainties.

Region	DJF	MAM	JJA	SON
Davis	54.5 (102%)	47.7 ( 97%)	30.7 (114%)	18.8 (127%)
Macquarie Island	61.3 ( 85%)	70.7 ( 91%)	17.9 (139%)	7.7 (229%)
Melbourne	96.7 (103%)	88.6 ( 89%)	26.7 (102%)	21.4 (109%)

#### 4.6.2 Results

The top panel of Fig. 4.12 shows the STT ozone enhancements, based on a vertical integration of the ozone above baseline levels for each ozonesonde where an event was detected. The area considered to be ‘enhanced’ ozone is outlined with yellow dashes on the left panel of Fig. 4.4. We find that the mean ozone flux associated with STT events is  $\sim 0.5\text{--}2.0 \times 10^{16} \text{ molecules cm}^{-2}$ . The bottom panel shows the mean fraction of total tropospheric column ozone (calculated from ozonesonde profiles) attributed to stratospheric ozone intrusions at each site for days when an STT event occurred. First the tropospheric ozone column is calculated, then the enhanced ozone column amount is used to determine the relative increase. At all sites, the mean fraction of tropospheric ozone attributed to STT events is  $\sim 1.0\text{--}3.5\%$ . On three separate days over Macquarie and Melbourne, this value exceeds 10%.

The upper panels in figures 4.13–4.15 show the factors  $I$ ,  $P$ , and  $\Omega_{\text{O}_3}$  which are used along with the assumed event lifetime to estimate the STT flux. The tropospheric ozone and area of our region is calculated using the output and surface area from GEOS-Chem over our three regions. The lower panel of these figures show the results of the calculation when we choose two days for our flux estimation, with dotted lines showing the range of flux calculated if we assume events last from one day to one week. The seasonal cycle of ozone flux is mostly driven by the  $P$  term, which peaks in the SH summer over all three sites. Total uncertainty (shaded) is on the order of 100% (see Sect. 4.7.2). We calculate the annual amount based on the sum of the monthly values. The regions over Davis, Macquarie Island, and Melbourne have estimated STT ozone contributions of  $\sim 5.7 \times 10^{17}$ ,  $\sim 5.7 \times 10^{17}$ , and  $\sim 8.7 \times 10^{17}$  molecules  $\text{cm}^{-2} \text{ a}^{-1}$  respectively, or equivalently  $\sim 2.0$ ,  $2.1$ , and  $3.3 \text{ Tg a}^{-1}$ .

#### 4.6.3 Comparison to literature

Škerlak, Sprenger, and Wernli 2014 show an estimate of roughly  $40$  to  $150 \text{ kg km}^{-2} \text{ month}^{-1}$  in these regions, over all seasons (see Fig. 16, 17 in their publication) while we estimate from  $0$  to  $180 \text{ kg km}^{-2} \text{ month}^{-1}$  STT impact, following a seasonal cycle with the maximum in austral summer. We estimate higher maximum flux over Melbourne, ( $178$ , and  $150 \text{ kg km}^{-2} \text{ month}^{-1}$  in January and February) than in either Davis ( $89 \text{ kg km}^{-2} \text{ month}^{-1}$  in March) or Macquarie Island ( $68 \text{ kg km}^{-2} \text{ month}^{-1}$  in January). Our calculated seasonal contributions, along with total uncertainty are shown in Table 4.3.

This result disagrees with several other studies which have found STT ozone fluxes in the SH extra-tropics are largest from autumn or winter to early spring. Roelofs and Lelieveld 1997 used a model carrying a tracer for stratospheric ozone to estimate STT

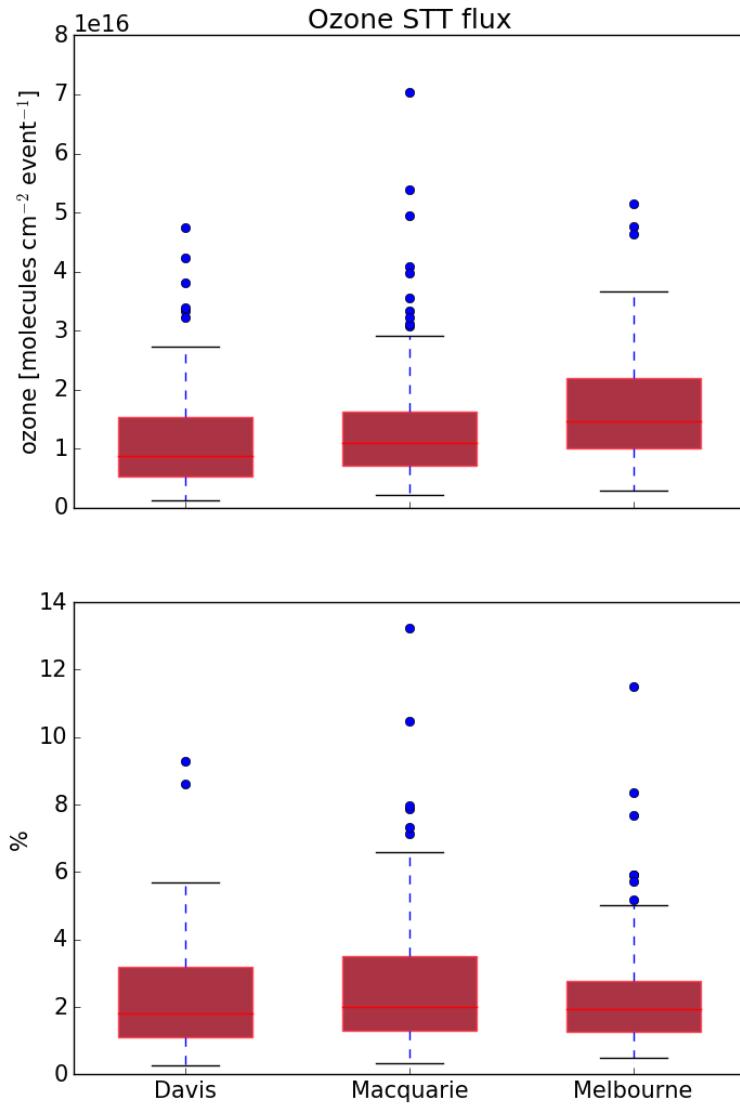


FIGURE 4.12: Top panel: tropospheric ozone attributed to STT events. Bottom panel: percent of total tropospheric column ozone attributed to STT events. Boxes show the inter-quartile range (IQR), with the centre line being the median, whiskers show the minimum and maximum, circles show values which lie more than 1.5 IQR from the median. Values calculated from ozonesonde measurements as described in the text.

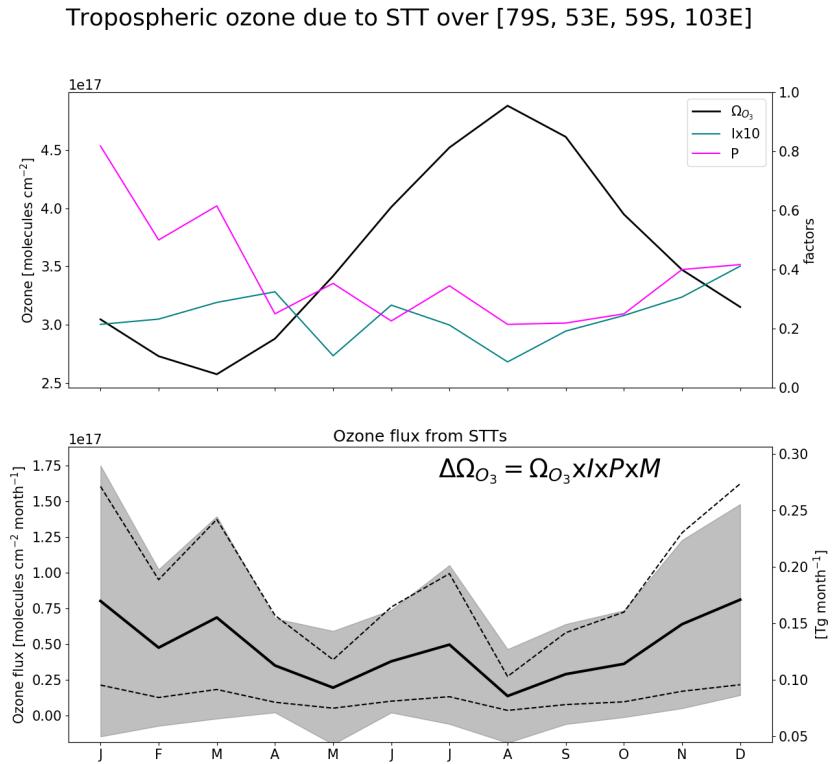


FIGURE 4.13: (Top) Tropospheric ozone, ( $I$ )mpact per event, and ( $P$ )robability of event detection per sonde launch, averaged over the region above Davis. The tropospheric ozone column  $\Omega_{O_3}$  (black, left axis) is from GEOS-Chem, while the STT probability  $P$ (magenta, right axis) and impact  $I$  (teal, right axis) are from the ozonesonde measurements. The STT impact is multiplied by ten to better show the seasonality. (Bottom) Estimated contribution of STT to tropospheric ozone columns over the region, with uncertainty (shaded area) estimated as outlined in Sect. 4.7. The black line shows STT ozone flux if event lifetime is assumed to be two days, with dashed lines showing the range of flux estimation if we assumed events lasted from one day to one week.

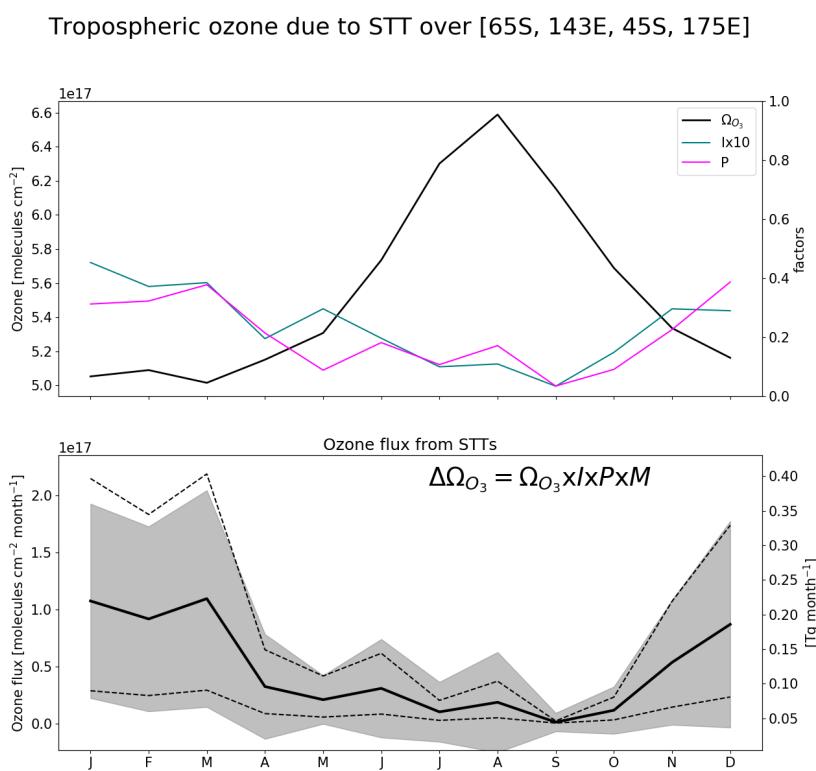


FIGURE 4.14: As described in 4.13, for the region containing Macquarie Island.

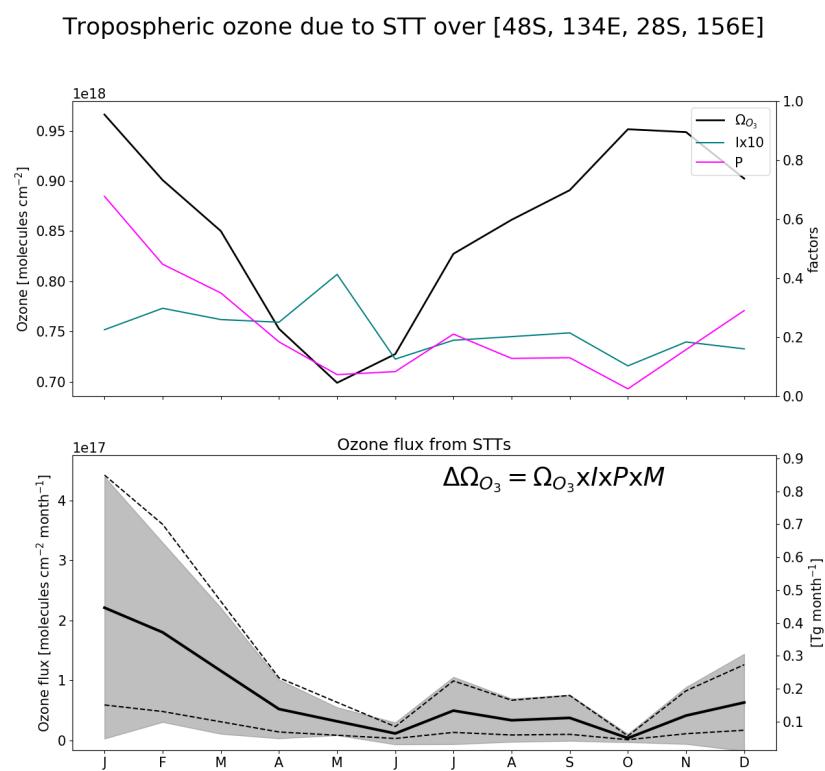


FIGURE 4.15: As described in 4.13, for the region containing Melbourne.

impacts. They see higher SH tropospheric ozone concentrations, as well as STT flux, in the SH winter. Our model also shows ozone column amounts peaking in winter, however flux is maximised in summer due to our detected event frequencies. Elbern, Hendricks, and Ebel 1998 examine STT using ECMWF data for prior to 1996, using PV and Q-vectors to determine STT frequency and strength, and suggest fewer fold events in the SH occur from December to February. Olsen (2003) used PV and winds from the GEOS reanalysis combined with ozone measurements from the TOMS satellite to estimate that the ozone flux between  $30^{\circ}$  S and  $60^{\circ}$  S is  $210 \text{ Tg yr}^{-1}$ , with the maximum occurring over SH winter. Liu et al. (2017a) model the upper tropospheric ozone and its source (emissions/lightning/stratospheric) over the Atlantic ocean between  $30^{\circ}$  S and  $45^{\circ}$  S, and suggest that most of this is transported from the stratosphere from March to September, which is when the subtropical jet system is strongest.

The disagreements largely reflect the difference between point source based estimates and zonally averaged estimates, as the meteorological behaviour at our three sites is not the same as the system that dominates the southern hemisphere in general. As detailed in Sect. 4.4, the maximum STT influx which occurs during SH winter is almost entirely due to the dominant STT system which occurs annually over the southern Indian ocean and middle of Australia. It is difficult to compare remote ozonesonde datasets with area averaged models or re-analyses based on non-co-located measurements (such as ERA).

## 4.7 Sensitivities and limitations

### 4.7.1 Event detection

Our method uses several subjectively-defined quantities in the process of STT event detection. Here we briefly discuss these quantities and the sensitivity of the method to each. Using the algorithm discussed in Sect. 4.3.3, we detect 80 events at Davis, 105 (21 fire influenced) events at Macquarie Island, and 127 (27 fire influenced) events at Melbourne.

The cut-off threshold (defined separately for each site) is determined from the 95th percentile of the ozone perturbation profiles between 2 km above the earth's surface and 1 km below the tropopause. We use the 95th percentile because at this point the filter locates clear events with fewer than 5% obvious false positive detections. Event detection is sensitive to this choice; for example, using the 96th, and 97th percentile instead decreased detected events by 2, 9 (2,10%) at Davis, 13, 31 (11, 28%) at Macquarie Island, and 8, 24 (6, 18%) at Melbourne. Event detection is therefore also sensitive to the range over which the percentile is calculated. This range was chosen to remove anomalous edge effects of the Fourier bandpass filter and to discount the highly variable ozone concentration which occurs near the tropopause.

Ozone enhancements are only considered STT events if they occur from 4 km altitude up to 500 m below the tropopause. This range removes possible ground pollution and events not sufficiently separated from the stratosphere, while still capturing many well-defined events that occur within 1 km of the tropopause. An example of a well-defined event that occurs within 1 km of the tropopause is shown in the supplementary (Fig. S2). However, STT events which reach below 4 km are physically

possible and we may have some false negative detections due to the altitude restricted detections.

### 4.7.2 Flux calculations

Flux is calculated as  $I \times P \times M \times \Omega_{O_3}$ , with each term calculated as described in Sect. 4.6.1. The uncertainty is determined using the standard deviation of the product, with variance calculated using the variance of a product formula, assuming that each of our terms is independent:

$$\text{var}(\Pi_i X_i) = \Pi_i (\text{var}(X_i) + E(X_i)^2) - (\Pi_i E(X_i))^2$$

The standard deviations for the  $I$  and  $\Omega_{O_3}$  terms are calculated over the entire dataset. These terms are considered to be homoskedastic (unchanging variance over time). Uncertainty in assumed event lifetime is set at 50%, as we believe it is reasonable to expect events to last 1-3 days.  $P$  is the probability of any ozonesonde detecting an event, and is assumed to be constant (for any month). The overall uncertainty as a percentage is shown in parentheses in Table 4.3, these values are on the order of 100%, largely due to relative uncertainty in the  $I$  factor which ranges from 50-120% for each month.

Small changes in the region don't have a large affect on the per area flux calculations: increasing or decreasing the regions by  $1^\circ$  on each side ( $\sim 10\%$  change in area) change the resulting flux by  $\sim 1\%$ . However due to the large portion of winter STT events being flagged due to potential smoke plume influence, a significant change in the yearly flux is seen when we don't remove these events. Without removing smoke flagged events we see an increase in estimated yearly flux of  $\sim 1.1, 2.1 \times 10^{17}$  molecules  $\text{cm}^{-2} \text{ yr}^{-1}$  (which is a change of  $\sim 15, 20\%$ ), over Macquarie Island and Melbourne respectively.

Considering the  $I$  factor, as discussed in here and in Sect. 4.7, there are several uncertainties in our method that are likely to lead to a low bias, such as the conservative estimate of flux within each event. Although there is little available data on SH ozone events for us to compare against, consider Terao et al. (2008), who estimated that up to 30–40% of the ozone at 500 hPa was transported from the stratosphere, in the northern hemisphere.

Our STT event impact estimates have some sensitivity to our biomass burning filter: including smoke-influenced days increases the mean per area flux by 15-20%. Although events which are detected near fire smoke plumes are removed, some portion of these could be actual STTs. The change in our  $P$  parameter when we include potentially smoke influenced events leads to a yearly estimated STT of  $11 \times 10^{17}$  molecules  $\text{cm}^{-2} \text{ yr}^{-1}$  over Melbourne, which suggests that up to  $2.1 \times 10^{17}$  molecules  $\text{cm}^{-2} \text{ yr}^{-1}$  ozone enhancement could be caused by smoke plume transported precursors. This is a potential area for improvement, as a better method of determining smoke influenced columns would improve confidence in our estimate.

Other possibly important uncertainties in our calculation of STT flux which we don't cover are listed here. Filtering events which occur within 500 m of the tropopause may also lead to more false negatives. This could also cause lower impact estimates due to only measuring ozone enhancements which have descended and potentially

slightly dissipated. On the other hand we have no measure of how often the detached ozone intrusion reascends into the stratosphere, which would lead to a reduced stratospheric impact. The estimated tropospheric ozone columns modelled by GEOS-Chem may be biased, for instance Hu et al. 2017 suggest that in general GEOS-Chem (with GEOS-5 met. fields) underestimates STT, with  $\sim 360 \text{ Tg a}^{-1}$  simulated globally, compared to  $\sim 550 \text{ Tg a}^{-1}$  observationally constrained. Transport uncertainty is very difficult to estimate with the disparate point measurements; it's possible that detected events are (at least partially) advected out of the analysis regions, which would mean we overestimate the influx into the region, and it is also possible that we are influenced by STT events outside the regions of analysis. Uncertainty in event longevity is set to 50%, however this implies a very simplistic model of event lifetimes. A great deal of work could be done to properly model the regional event lifetimes, however this is beyond the scope of our work.

Uncertainties in STT ozone flux detection are ( $\sim 100\%$ ), and could be directly improved with larger or longer datasets. Possible parameterisations and an improved model of event lifetime could also improve the confidence in our estimate of event impacts, as well as allowing fewer assumptions.

## 4.8 Conclusions

Stratosphere-to-troposphere transport (STT) can be a major source of ozone to the remote free troposphere, but the occurrence and influence of STT events remains poorly quantified in the southern extra-tropics. Ozonesonde observations in the SH provide a satellite-independent quantification of the frequency of STT events, as well as an estimate of their impact and source. Using almost ten years of ozonesonde profiles over the southern high latitudes, we have quantified the frequency, seasonality, and altitude distributions of STT events in the SH extra-tropics. By combining this information with ozone column estimates from a global chemical transport model, we provided a first, conservative estimate of the influence of STT events on tropospheric ozone over the Southern Ocean.

Our method involved applying a bandpass filter to the measured ozone profiles to determine STT event occurrence and strength. The filter removed seasonal influences and allowed clear detection of ozone-enhanced tongues of air in the troposphere. By setting empirically-derived thresholds, this method clearly distinguished tropospheric ozone enhancements that are separated from the stratosphere. Our method is sensitive to various parameters involved in the calculation; however, for our sites we saw few false positive detections of STT events.

Detected STT events at three sites spanning the SH extra-tropics ( $38^\circ\text{S}$ ,  $55^\circ\text{S}$ , and  $69^\circ\text{S}$ ) showed a distinct seasonal cycle. All three sites displayed a summer (DJF) maximum and an autumn to winter (AMJJA) minimum, although the seasonal amplitude was less apparent at the Antarctic site (Davis) as events were also detected regularly in winter and spring (likely due to polar jet stream-caused turbulence). Analysis of ERA-Interim reanalysis data suggested the majority of events were caused by turbulent weather in the upper troposphere due to low pressure fronts, followed by cut-off low pressure systems. Comparison of ozonesonde-measured ozone profiles against those simulated by the GEOS-Chem global chemical transport model showed the model is

able to reproduce seasonal features but does not have sufficient vertical resolution to distinguish STT events.

By combining the simulated tropospheric column ozone from GEOS-Chem with ozonesonde-derived STT estimates, we provide a first estimate of the total contribution of STT events to tropospheric ozone in these southern extra-tropical regions. We estimate that the ozone enhancement due to STT events near our three sites ranges from 300–570 kg km<sup>-2</sup> a<sup>-1</sup>, with seasonal maximum in SH summer.

Estimating STT flux using ozonesonde data alone remains challenging; however, the very high vertical resolution provided by ozonesondes suggests they are capable of detecting STT events that models, re-analyses, and satellites may not. Further work is needed to more accurately translate these ozonesonde measurements into STT ozone fluxes, particularly in the SH where data are sparse and STT is likely to be a major contributor to upper tropospheric ozone in some regions. More frequent ozonesonde releases at SH sites could facilitate development of better STT flux estimates for this region.

## 4.9 Contributions and Acknowledgements

JWG wrote the algorithms, ran the GEOS-Chem simulations, performed the analysis and led the writing of the paper under the supervision and guidance of SPA, RS, and JAF. AK contributed the Davis ozonesonde data and performed the analysis of the alternate STT proxy. All authors contributed to editing and revising the manuscript.

*Data availability.* All GEOS-Chem model output and the ozonesonde observational data are available from the authors upon request.

We thank Dr. Sandy Burden for help clarifying some of the uncertainties involved in methods within this work. We also thank Dr. Clare Paton-Walsh, who identified the need to account for smoke-influenced events, and provided discussions on how to go about doing such. Ozonesonde data comes from the World Ozone and Ultraviolet Data Centre (WOUDC). The ERA-Interim data were downloaded from the ECMWF website following registration. This research was undertaken with the assistance of resources provided at the NCI National Facility systems at the Australian National University through the National Computational Merit Allocation Scheme supported by the Australian Government. This work was supported through funding by the Australian Government's Australian Antarctic science grant program (FoRCES 4012), the Australian Research Council's Centre of Excellence for Climate System Science (CE110001028), the Commonwealth Department of the Environment ozone summer scholar program. This research is supported by an Australian Government Research Training Program (RTP) Scholarship.



## Chapter 5

# Summary and Concluding Remarks

**In this thesis I aimed to improve understanding of natural contributions to ozone over Australia and the southern ocean.** Chapter 1 Introduction and Literature Review examined processes leading to ozone enhancement in the troposphere, primarily isoprene emissions and subsequent chemistry. A secondary source of ozone, stratosphere to troposphere transport (STT), was also outlined with associated processes and causes discussed. A summary of how the lack of information available over Australia affects modelling and forecasting was provided. Methodologies, tools and data-sets used throughout the thesis were detailed in Chapter 2 Data and Modelling . Chapter 3 Biogenic Isoprene Emissions in Australia showed how models are misrepresenting isoprene emissions by a large margin in Australia, along with the description and implementation of a relatively simple method of improvement and its impacts and uncertainties. Chapter 4 Stratospheric ozone intrusions provided an estimate of stratospheric ozone influx to the troposphere, along with potential classifications and seasonality.

### 5.1 Outcomes

**I aimed to recalculate satellite vertical columns of HCHO using updated model a priori information.** HCHO from the OMI instrument onboard the Aura satellite was examined and recalculated in Chapter 2, where the influence of the air mass factor (AMF) was discussed in detail. The original AMF used in the satellite product was created using an older version of the GEOS-Chem model with out-of-date HCHO chemistry. Recalculation, binning, and analysis of the satellite HCHO vertical columns was performed using GEOS-Chem v10.01, outlined in Section 2.6.1. This was performed using [newly developed my own<sup>mh</sup>](#) partial AMF recalculation code, along with full recalculation code set up in collaboration with Prof. Palmer and Dr. Surl. Satellite measured vertical columns changed systematically when using updated shape factors, and again when using updated shape factors in conjunction with updated scattering weights. Over Australia an increase in average total column HCHO was observed, along with lower background levels, following either recalculation. Even though complete recalculation is computationally more expensive, it is recommended as the differences seen with recalculated scattering weights are substantial.

**I aimed to determine biogenic isoprene emissions in Australia using a top-down inversion of satellite HCHO, through an estimated yield from GEOS-Chem.** In Chapter 3 [I determined<sup>mh</sup>](#) the linear relationship between total column HCHO and

biogenic isoprene emissions over Australia is determined<sup>mh</sup> using the global model GEOS-Chem. Applying this relationship to satellite HCHO measurements created the desired top-down isoprene emissions estimate. This process was described in Section 3.2, using extensive filtering of both satellite data (to exclude non-biogenic HCHO sources) and model yield (to minimise spatial smearing). The uncertainty and limitations of top-down estimates due to satellite and model uncertainty, along with temporal and horizontal resolution of available data, were examined in Section 3.4. Furthermore, this top-down estimation was used to scale isoprene emissions for a new simulation of HCHO and O<sub>3</sub> over Australia, described in Section 3.2.9. MEGAN isoprene emissions were shown to be overestimated in summer by a factor of 2-5, with total emissions of 39 Tg yr<sup>-1</sup> in the a priori dropping to 21 Tg yr<sup>-1</sup> in the a posteriori. Reduced isoprene emissions across Australia was shown to lower ozone concentrations by approximately 5% mostly in the surface level of model output.

**I aimed to improve understanding of ozone transported to the troposphere from the stratosphere over Australia and the southern ocean.** In Chapter 4 the seasonal cycle of STT events was characterised, and their contribution to the SH extra-tropical tropospheric ozone budget quantified using GEOS-Chem to estimate ozone flux extrapolated from three measurement sites. Causal climatology and event seasonality were examined in Section 4.4. STT detection frequencies and modelled tropospheric ozone columns were used to estimate STT ozone flux near three sites in Section 4.6. Findings were compared against relevant literature, and the uncertainties involved in STT event detection and ozone flux estimation have also been examined.

## 5.2 Isoprene emissions

A new isoprene emissions estimate was created using OMI satellite measurements of HCHO. This estimate was compared against the bottom-up estimate from GEOS-Chem (running MEGAN). Generally months outside of May to August showed the a posteriori lower than the a priori, except in the southeastern portion of Australia where only February emissions were notably reduced. MEGAN isoprene emissions appear to be overestimated in austral summer, suggesting poorly understood emission factors for Australian forests. The overall reduction over Australia was ~ 46%, and the MEGAN based a priori emissions range from 2-5 times higher than the a posteriori.

## 5.3 Tropospheric Ozone over Australia

Ozone production in the troposphere is a complex process involving various compounds. Tropospheric ozone is enhanced through biogenic VOC chemistry, stratospheric transport, and pollution. The first two of these processes are highly uncertain, with few studies performed on either topic within Australia. Of these ozone sources, VOC chemistry uncertainty is dominated by poor understanding of biogenic emissions (which are mainly isoprene). Emissions of isoprene have been globally modelled at ~465-500 Tg C yr<sup>-1</sup> (Guenther et al. 2006; Messina et al. 2016). However these appear to be overestimated in Australia, as seen here with top-down estimates at ~ 18

Tg C yr<sup>-1</sup> compared to ~ 35 Tg C yr<sup>-1</sup> from bottom-up estimates using GEOS-Chem (which implements MEGAN).

Tropospheric ozone production was estimated using GEOS-Chem before and after scaling isoprene emissions based on top-down satellite data. This was summarised in Figure 3.31 showing summer surface ozone differences before and after scaling isoprene emissions. Figure 5.1 shows Australian summer surface level ozone with and without scaling isoprene emissions to match the multi-year top-down estimation.<sup>tb</sup> The reduction in surface ozone is approximately 5%. Total column ozone does not change much (<< 1%) and most of the change in ozone is only seen at the surface level. MThis shows that modelled impacts from reduced isoprene emission are seen nearly completely in the surface level, and outweighed by other processes at higher altitudes.

**Figure 5.1 deleted from this section.<sup>tb</sup>**

Ozone is one of the two measured air pollutants that have not fallen in concentration over the last 10 years (Keywood, Emmerson, and Hibberd 2016). Isoprene emissions are one of the important precursors to ozone production, and this has been examined in Section 3.3. The results are tabulated for Australia and five sub-regions, in Table 3.7. The sub-regions were detailed in Section 3.3 and shown again here in Figure 5.2<sup>tb</sup>. A 5% mean difference in surface ozone levels will directly affect population exposure levels and impacts air quality estimates. This effect could be underestimated in cities due to the low horizontal resolution used in this thesis, as isoprene is most likely to impact city fringes (Millet et al. 2016). The background ozone changes seen here need to be accurately modelled to confidently predict ozone exceedances (e.g., Cope et al. 2004).

**Table 5.1 deleted from this section.<sup>tb</sup>**

**Figure 5.2 deleted from this section.<sup>tb</sup>**

Ozone transported from the stratosphere affects surface ozone levels, although this was hard to detect with the methods used in this thesis. The effects of two influences on the tropospheric ozone column were examined in this thesis. STT analysis over Melbourne suggests up to ~ 10% of the tropospheric column was due to stratospheric influx, with the average increase caused by STT influx being ~ 1 to 3.5%. From the biogenic source a relatively large (~ 50%) drop in isoprene emissions causes a ~ 1 – 5% reduction in surface ozone concentrations over summer. The direct correlation between reduced emissions and surface ozone reductions was not apparent without taking large area averages, likely due to the effects being downwind of where isoprene emissions are reduced. Both STT and biogenic ozone sources become more important in summer, as STT occur more frequently and isoprene emissions increase with the summer high temperatures. Days with exceptionally high temperatures can also lead to high ozone concentrations not shown in models (e.g., Paton-Walsh et al. 2018). The impacts of these two sources vary with altitude with most STT occurring in the free troposphere, while isoprene emission impacts were largely seen at the surface. Ozone in the free troposphere has the largest effect on climate, while surface ozone matters more for air quality and ecosystem impacts. Therefore, accurate STT modelling is expected to impact estimates of radiative forcing, while improved isoprene emission modelling is expected to improve air quality and surface ozone concentration simulations.

## 5.4 Outputs

Analysis of STT along with an estimate of ozone flux from the stratosphere over parts of Australia and the southern ocean was published [as in ACP](#): Jesse W Greenslade et al. (2017). "Stratospheric ozone intrusion events and their impacts on tropospheric ozone in the Southern Hemisphere". In: *Atmospheric Chemistry and Physics*, pp. 1–33. URL: <https://www.atmos-chem-phys.net/17/10269/2017/acp-17-10269-2017.pdf>.

A top-down estimate of isoprene emissions over Australia created using OMI HCHO and GEOS-Chem modelled chemistry in Chapter 3 is currently in preparation: Greenslade J. W., Fisher J. A. [et al.](#), Top-down emissions of isoprene over Australia, in prep. Re-gridded OMI HCHO prepared in this thesis has been used in a recent publication (of which I am a co-author), which examines the HCHO trend over Wollongong over the last 20 years: Kaitlyn J. Lieschke et al. (2019). "Decreasing trend in formaldehyde detected from 20-year record at Wollongong, Southeast Australia". In: *Geophysical Research Letters*. DOI: 10.1029/2019GL083757. URL: <https://doi.org/10.1029/2019GL083757>.

## 5.5 Future work

This section briefly examines possible future works or extensions from this thesis that would be beneficial to tropospheric ozone (and precursor) modelling over Australia.

One important parameter potentially influencing isoprene emission in Australia is the soil moisture activity factor ( $\gamma_{SM}$ ) (Sindelarova et al. 2014; Bauwens et al. 2016). Sindelarova et al. (2014) and more recently Emmerson et al. (2019) showed how Australian isoprene emissions could be as much as halved by accounting for lower soil moisture. Generally if soil moisture is too low, isoprene emissions stop (Pegoraro et al. 2004; Niinemets et al. 2010), however in many Australian regions the plants may be more adapted to lower moisture levels (Emmerson et al. 2019). These moisture states are not implemented in recent GEOS-Chem versions (including v10.01, used in this thesis). Improved isoprene emissions modelling requires this soil moisture problem to be addressed. Simply enabling the parameter in its current form is not sufficient for Australia, due to both the unknown soil moisture and the poorly understood plant responses in this country. Recently, updated soil moisture parameterisation has been shown to improve modelled isoprene emissions in drought conditions (Jiang et al. 2018). Alternatively improving the soil moisture and drought process parameterisation in GEOS-Chem could be performed following a similar process to that shown in Emmerson et al. (2019), using satellite based soil moisture estimates. Calibrating and applying a soil moisture update to Australia would be a worthwhile future project. Testing improvements and sensitivity of modelled VOCs and ozone to a soil moisture update could provide useful information with which to update GEOS-Chem.

Using satellite data to improve isoprene emission estimates such as has been done in this thesis must be used to analyse model improvements, since fully independent measurements are lacking. Measurements of isoprene emissions in Australia would provide a valuable opportunity to verify improvements in emission estimates.

Emissions measurement would be valuable however they remain expensive and difficult, especially over the large sparse environment that makes up much of the Australian continent. Working on measurement priorities for potential campaigns is a worthwhile future project, due to the limitations of coverage such operations can reasonably provide. For instance ground- and aircraft-based VOC, NO<sub>x</sub>, HCHO, and ozone measurements would greatly improve the understanding of both satellite biases and model uncertainties over Australia. Additionally, measurements in the northern forested areas of Australia could be used to improve how models handle the monsoonal conditions which are not likely to be accurately captured by satellite measurements due to increased cloud coverage.

This thesis examined isoprene emissions and how they affect ozone concentrations at one resolution, using one satellite as the basis for a top-down inversion. Other satellites (e.g., GOME-2) could be included to allow more robust estimates as well as providing some diurnal resolution as each satellite has a different overpass time. An adjoint version of GEOS-Chem could be implemented over a smaller domain (either spatially or temporally or both) to provide more detailed analysis of both ozone and isoprene transport. This sort of modelling would also provide understanding of the resolution limitations of top-down emissions estimates, along with a clearer look at how sensitive the technique is to various parameters.

Considering ozone flux estimates based on ozonesondes, the process outlined in Chapter 4 could be applied to other mid to high southern latitude ozonesondes to provide hemispheric-scale understanding of STT and its influences. The work as published could be improved to automatically filter potential fire smoke influence using freely available satellite data, which would lead to a very simple and computationally inexpensive method for STT detection. A similar improvement based on online (or user specified) ozone concentration data would allow quick estimation of STT impacts, which are highly uncertain in many areas (especially in the southern hemisphere).

## 5.6 Concluding remark

Overall this thesis has contributed to the understanding of two important natural sources of ozone over Australia by improving *theour<sup>mh</sup>* understanding of isoprene emissions and by improving the characterisation of stratospheric intrusions. Through chemical modelling, impacts on tropospheric ozone from these sources has been quantified and analysed for several regions over Australia. A top-down estimate of isoprene emissions, a dominant biogenic VOC and one of the least characterised in Australia, has also been created and analysed.



## Appendix A

# Relationship between temperature and HCHO

Biogenic HCHO concentrations are correlated with temperature, as isoprene emissions are strongly correlated with temperature (Palmer et al. 2006; Zhu et al. 2013; Surl, Palmer, and Abad 2018). Fires emit HCHO precursors and increase HCHO concentrations independently of the relationship between temperature and HCHO, and should be revealed as outliers when comparing HCHO to temperature. Figures A.1 - A.3 show the relationship between modelled temperature, and satellite HCHO for January 2005 within subsets of Australia. A reduced major axis regression is used to determine the correlation between surface temperature (X axis) and HCHO (Y axis). Using the natural log of HCHO we can take the linear regression and then exponentiate each side in the equation  $\ln Y = mX + b$  to get  $Y = \exp mX + b$ . This gives us the exponential fit as shown, with the correlation coefficient between  $\ln HCHO$  and temperature. The distributions of exponential correlation coefficient and  $m$  terms is shown in the embedded plot, with one data point available for each grid square where the regression is performed. These figures show that the modelled temperature is not well correlated with corrected recalculated OMI vertical columns ( $r$  ranges from -0.24 to 0.47), but is with modelled columns ( $r$  ranges from 0.49 to 0.84). Correlations between modelled temperatures and HCHO are further improved when using the spatial average within each region. Furthermore the relationship is improved in individual grid squares over south Eastern Australia by removing non-biogenic emissions from the model ( $r$  increased from 0.58 to 0.75). This improvement is not seen in Northern Australia, nor south Western Australia. Overall this suggests that modelled correlations between temperature and HCHO are spatially dependent, and generally well reflected in satellite measurements over Australia.

One problem with detecting outliers in the temperature and enhanced HCHO relationship is that days when fires occur are likely to be hot. Another problem with correlating heat and HCHO is that increased temperature accelerates HCHO destruction (Zheng et al. 2015). We test the fire mask (see Section 3.2.3.1) by examining the relationship between modelled temperature and satellite HCHO with and without applying the filters for smoke and active fires. Figure A.4 show the regressions between mid-day surface HCHO and temperature, where two independent data sets are used for the temperature: one from GEOS-Chem output and the other from CPC daily maxima. Regressions over both the grid square containing Sydney and averaged over a wider area are reduced in quality when after applying the pyrogenic filter. This could be

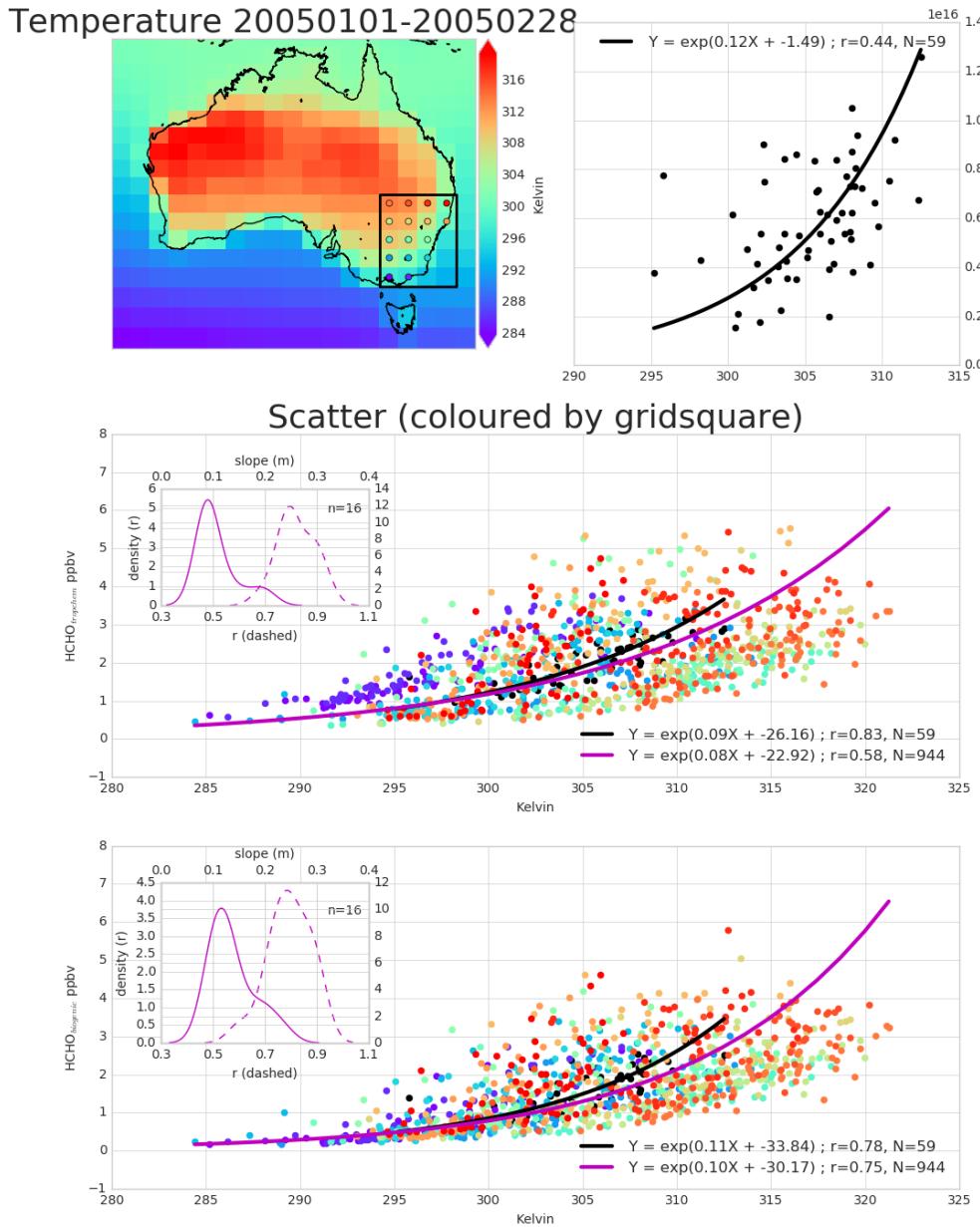


FIGURE A.1: Top row (left): surface temperature averaged over January and February 2005. Top row (right): correlation between spatially averaged GEOS-Chem temperatures and recalculated satellite vertical columns. Second row: GEOS-Chem surface temperatures correlated against GEOS-Chem HCHO, with different colours for each grid box, and black showing the spatially averaged correlation over time. Third row: as second row, except GEOS-Chem HCHO comes from the bio-genic emissions only simulation. A reduced major axis regression is used within each gridbox using daily overpass time surface temperature and HCHO. The distribution of slopes (solid) and regression correlation coefficients (dashed) for the exponential regressions is shown in the inset panels in rows 2 and 3.

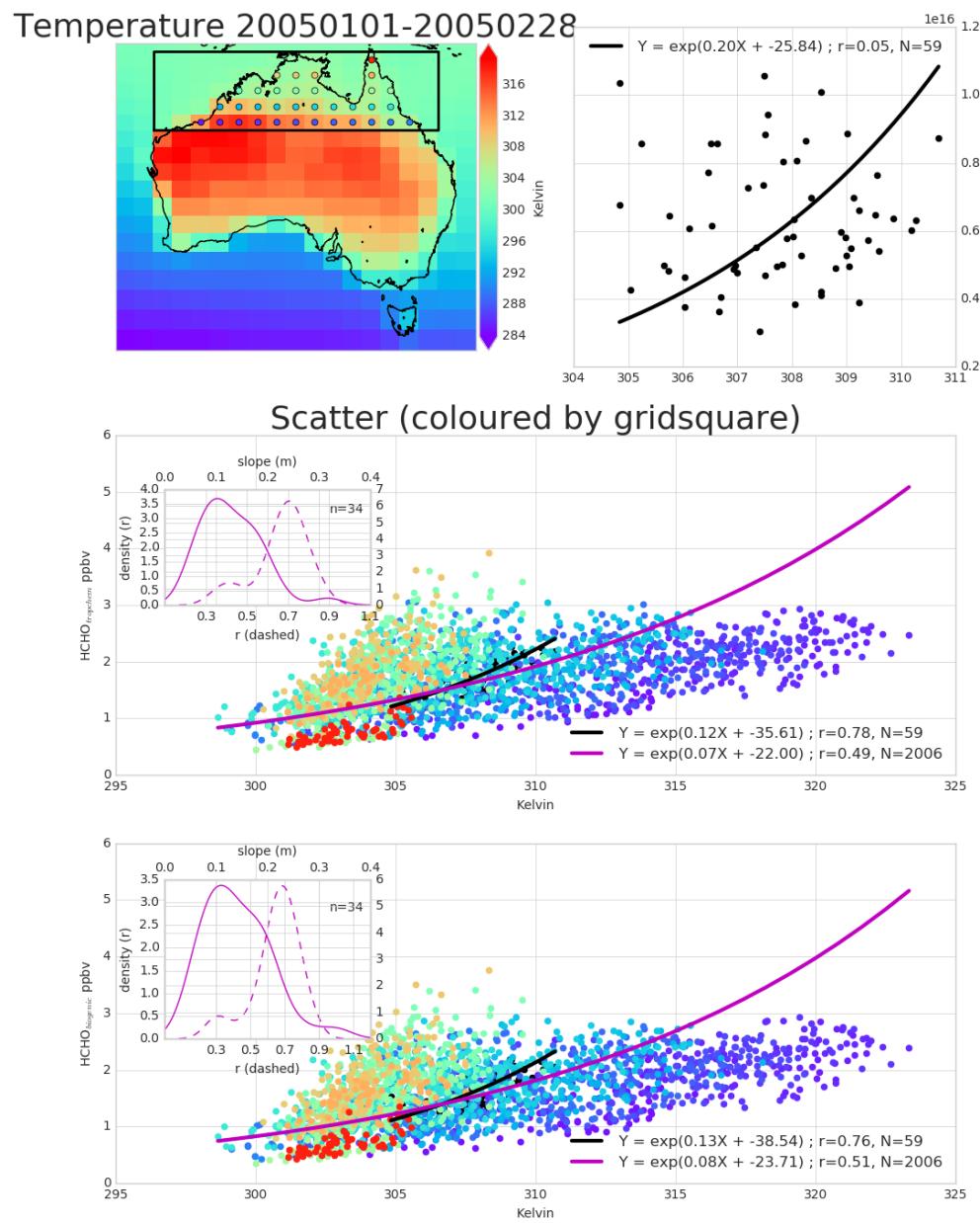


FIGURE A.2: As Figure A.1 but for northern Australia.

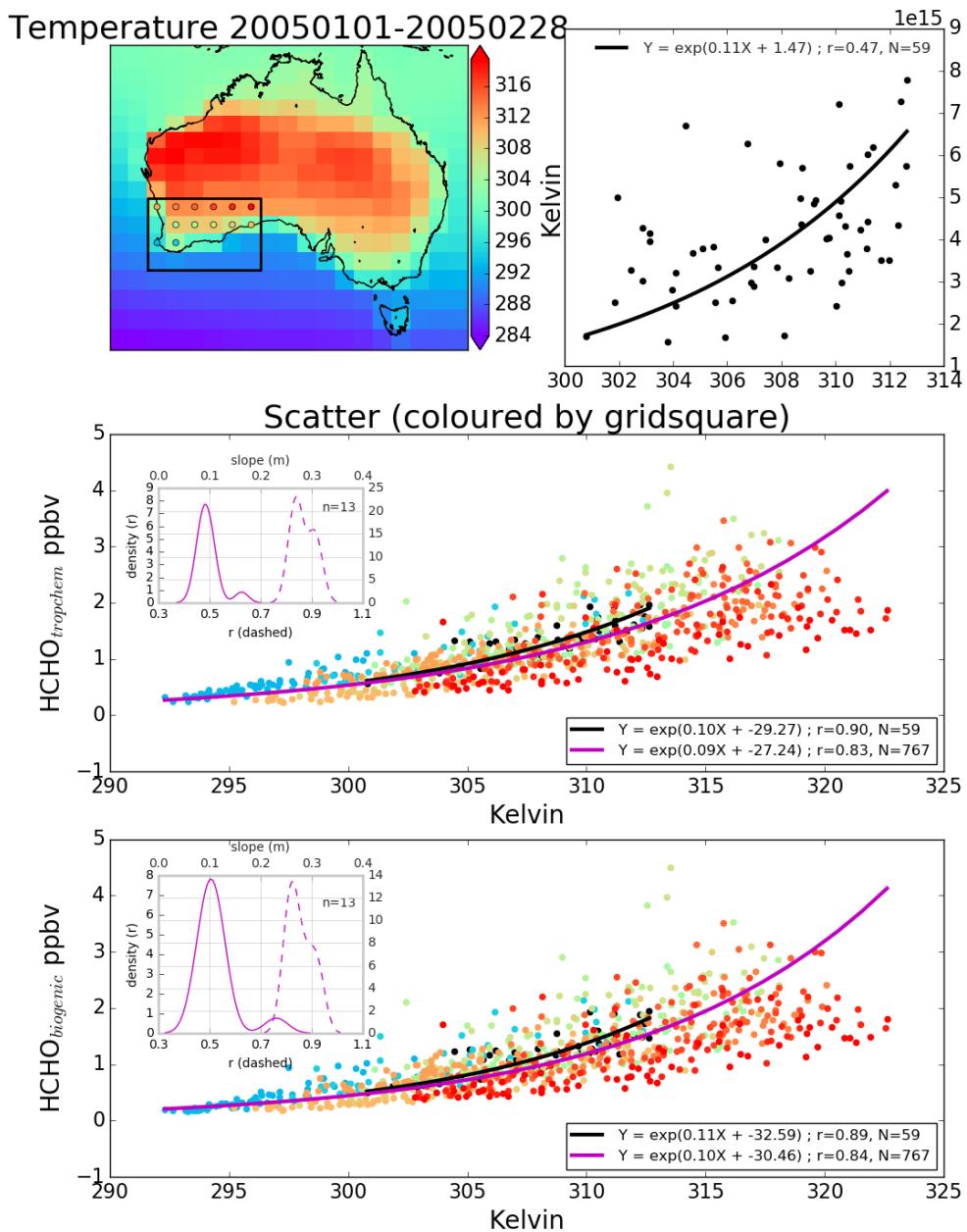


FIGURE A.3: As Figure A.1 but for south-western Australia. [TODO: Fix y axis in subplot 322](#)

due to the low resolution of available GEOS-Chem midday output, which greatly increases the strictness of the fire filter, which originally is created at  $0.25^\circ \times 0.3125^\circ$  horizontal resolution but here is widened out to  $2^\circ \times 2.5^\circ$ . While the modelled correlation between surface HCHO and temperature is quite strong ( $r > 0.8$ ) the correlation coefficient is reduced while the slope (temperature coefficient) is changed by more than 30% when applying the pyrogenic filter at the model resolution. If we use recalculated satellite HCHO columns instead of modelled midday outputs, then higher horizontal resolution can be achieved; however, this is at the cost of vertical resolution. Figure A.5 shows the relationship between total column HCHO and temperature, which is not as strong as that shown between surface HCHO and temperature. At this resolution the application of the pyrogenic filter is shown to slightly strengthen the correlation over the wider south eastern Australian region, with  $r$  increasing from  $\sim 0.31$  to  $\sim 0.38$ . In this thesis the pyrogenic filter is applied at the higher horizontal resolution ( $0.25^\circ \times 0.3125^\circ$ ), and this analysis suggests that the filter should strengthen the relationship between total column HCHO and its biogenic precursors over Australia.

## Temperature vs surface HCHO [ppbv] 20051201-20060228

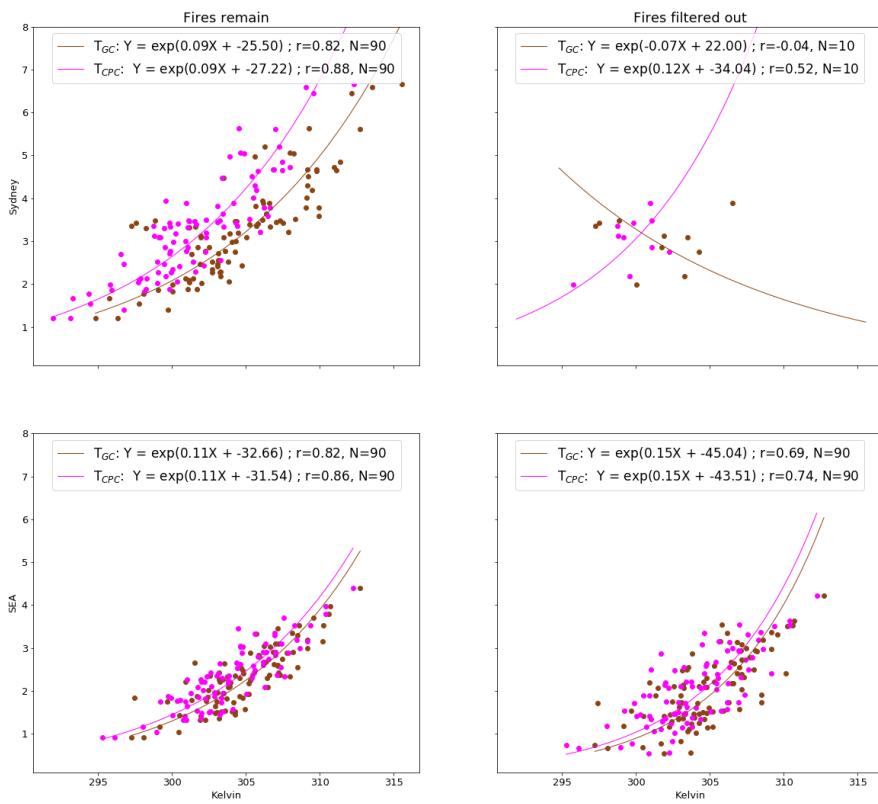


FIGURE A.4: Surface HCHO from GEOS-Chem overpass output (mid-day) on the Y axis vs surface temperatures at midday ( $T_{GC}$ ) and vs maximum daily temperatures from the CPC data set ( $T_{CPC}$ ). Top row: Sydney grid square scatter plot and regression with one data point for each day in the summer of 2005-2006. Bottom row: as top row except averaging over several grid boxes covering south eastern Australia (SEA: 37°S to 29°S, 146°E to 153.5°W). Grid squares with pyrogenic influence detected are removed (prior to any averaging) in the right column.

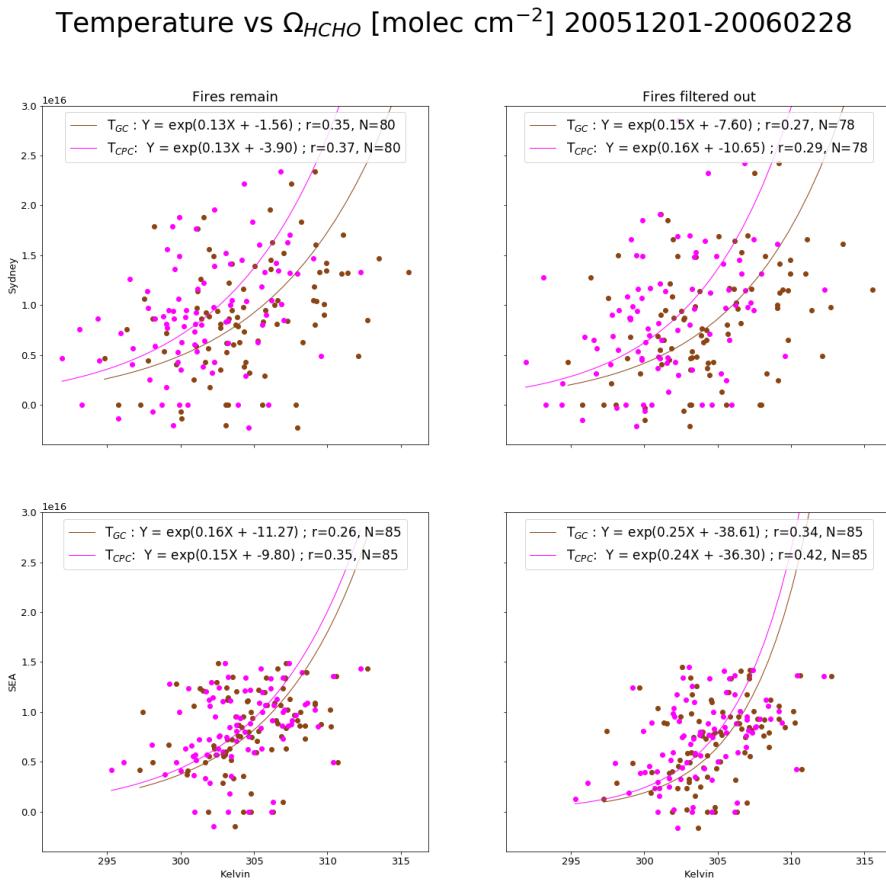


FIGURE A.5: OMI recalculated vertical columns of HCHO on the Y axis vs surface temperatures at midday ( $T_{GC}$ ) and vs maximum daily temperatures from the CPC data set ( $T_{CPC}$ ). Top row: Sydney grid square scatter plot and regression with one data point for each day in the summer of 2005-2006. Bottom row: as top row except averaging over several grid boxes covering south eastern Australia (SEA: 37°S to 29°S, 146°E to 153.5°W). Grid squares with pyrogenic influence detected are removed (prior to any averaging) in the right column.



## Appendix B

# Daily a priori regression against a posteriori

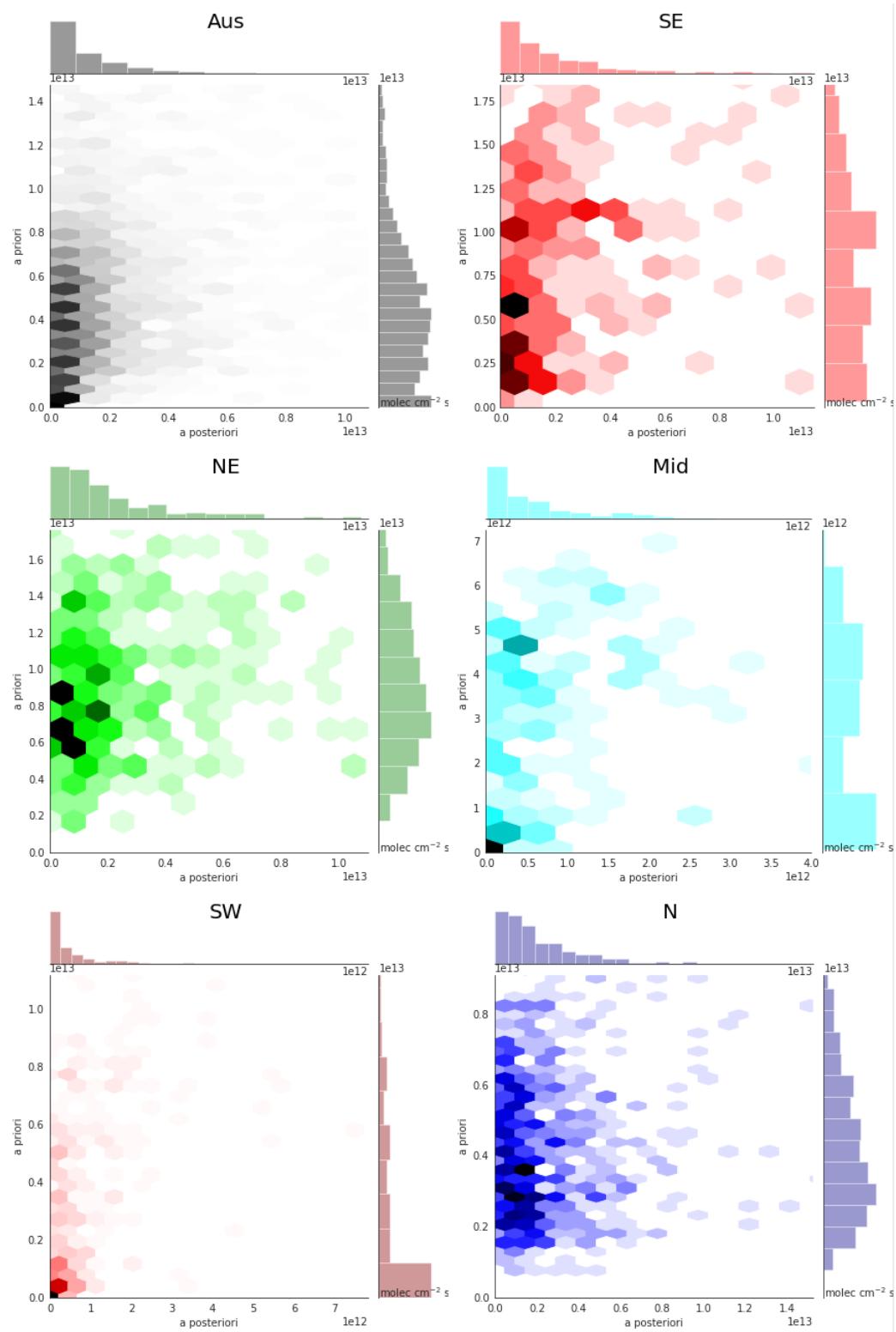


FIGURE B.1: Scatter plot (binned hexagonally with darker colours representing higher data-point frequency) along with the distributions of MEGAN (y axis) and the top-down estimate (x axis). This figure is based on summer (December - February) midday values over multiple years (2005-2012). Coloured by regions shown in Figure 3.17.

## Appendix C

# Supplementary for Ozone STT chapter

### S1 Foreword

This supplementary is reproduced as published (Jesse W Greenslade et al. (2017). "Stratospheric ozone intrusion events and their impacts on tropospheric ozone in the Southern Hemisphere". In: *Atmospheric Chemistry and Physics*, pp. 1–33. URL: [https://www.atmos-chem-phys.net/17/10269/2017.pdf](https://www.atmos-chem-phys.net/17/10269/2017/acp-17-10269-2017.pdf)) and available at <https://www.atmos-chem-phys.net/17/10269/2017/>.

### S2 Weather and Smoke analysis

Figure S1 contrasts two days; one with and one without signs of biomass burning influence near the Melbourne site (purple circle). On 17 October 2007 (top) elevated CO suggests the site may have been influenced by long-range transport from African and/or South American biomass burning. In contrast, on 3 February 2006 (bottom) CO columns across the SH show no influence from biomass burning.

Figure S2 (left) shows the vertical ozone profile on 3 February 2005. The tropopause was between 400 and 500 hPa and ozone in the upper troposphere was anticorrelated with relative humidity, suggesting the ozone enhancements are due to dry stratospheric air. An ozone intrusion into the troposphere at  $\sim$ 520 hPa was identified by our detection algorithm. The right panel shows the concurrent synoptic weather system, a cut-off low pressure system that caused a large storm and lowered the local tropopause height for several days. The flux of stratospheric ozone into the troposphere associated with this event, calculated using the method shown in Sect. 2.3 of the parent document, was at least  $3.1 \times 10^{11}$  molecules cm $^{-3}$ , or 8% of the tropospheric ozone column.

Figure S3 (left) shows the ozone profile over Melbourne on 13 January 2010. The tropopause was higher on this date (120-160 hPa). Using our algorithm, we detected an ozone intrusion centred around 200 hPa. As before, ozone anti-correlation with relative humidity provides further evidence that the elevated ozone was stratospheric in origin. In this profile, there was clear separation between the detected intrusion (highlighted in pink) and the ozone tropopause (black dashed line), which indicates that the sonde passed through regular tropospheric air after hitting a stratospheric intrusion

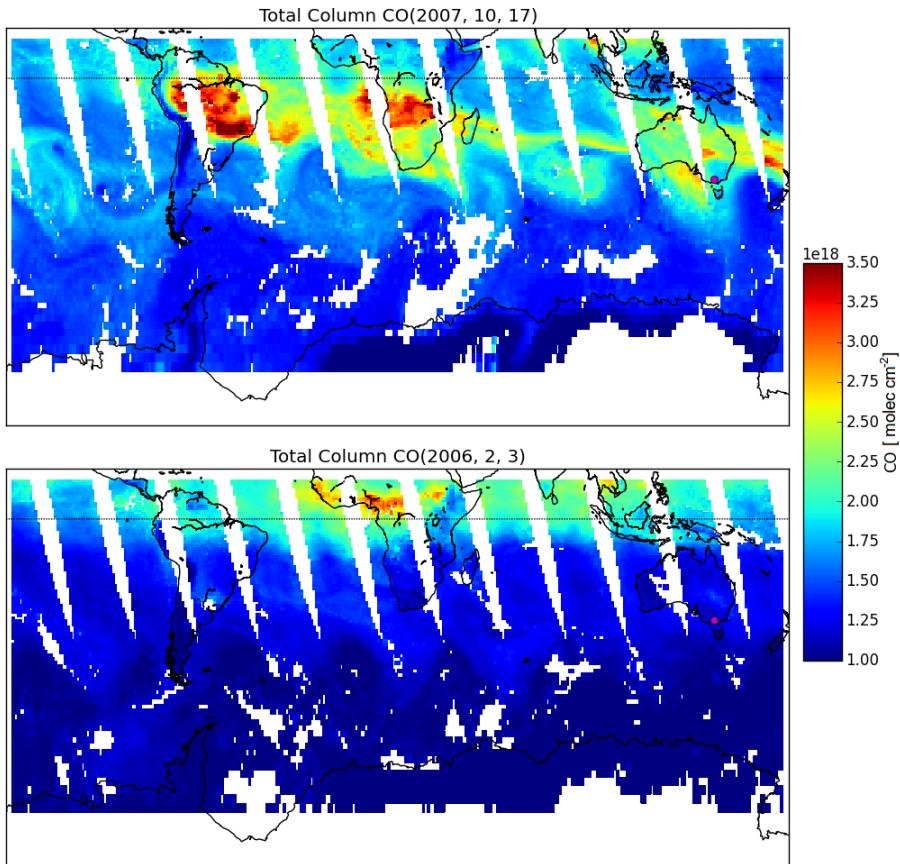


FIGURE S1: Example detection of biomass burning influence using AIRS total column CO. The top panel (17 October 2007) shows a day when ozone above Melbourne (purple dot) could have been caused by a transported biomass burning plume, and so was flagged in subsequent analysis. The bottom panel (3 February 2006) shows a day when ozone over Melbourne was not influenced by transported smoke.

but before reaching the tropopause. The right panel shows that this event was associated with a trough (front) of low pressure passing over south eastern Australia. This front travelled from west to east and caused a wave of lowered tropopause height. Frontal passage is a known cause of STT as stratospheric air descends and streamers of ozone-rich air break off and mix into the troposphere (Sprenger, Croci Maspoli, and Wernli 2003).

### S3 Southern Ocean extrapolation

#### S3.1 Outline

We use simulated tropospheric ozone columns from GEOS-Chem to extrapolate the ozonesonde-based estimates over a large area of the Southern Ocean encompassing

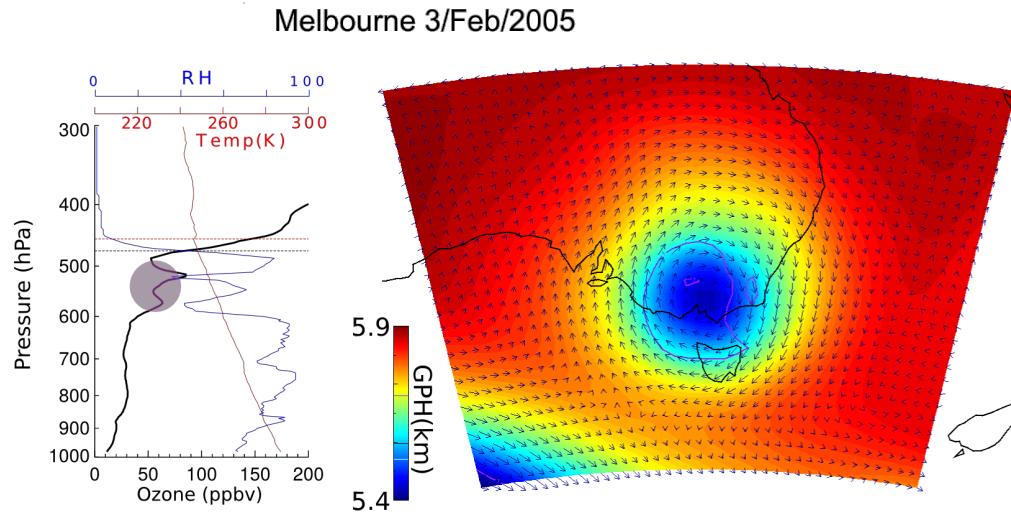


FIGURE S2: (Left) Vertical profile of ozone (black), relative humidity (blue), and temperature (red) measured by ozonesonde over Melbourne on 3 February 2005. The detected ozone STT event is highlighted in pink. Tropopause heights using both the ozone definition (black dashed line) and lapse rate definition (red dashed line) are also shown. (Right) Geopotential heights at 500 hPa from the ERA-Interim reanalysis, with wind vectors over-plotted. Also shown is the 1 PVU contour line (purple).

our three measurement sites. Figure S4 shows the region defined by latitudes  $79^{\circ}$  -  $28^{\circ}$  S, and longitudes  $53^{\circ}$  -  $175^{\circ}$  E.

Figure S5 (upper panel) shows the factors  $I$ ,  $P$ , and  $\Omega_{O_3}$  which are used along with the assumed event lifetime to estimate the STT flux. The tropospheric ozone and area of our region is calculated using the output and surface area from GEOS-Chem over the Southern Ocean grid boxes along with the molecules  $\text{cm}^{-2}$  per month calculations, along with ozone molar mass of  $48 \text{ g mol}^{-1}$ .

It is worth noting that this extrapolation is very simplistic and is performed as an example of how the seasonal ozone STT calculations could be used. A more spatially resolved estimate could be determined by dividing the Southern Ocean region into longitudinal and latitudinal bins for calculating the average  $\Omega_{O_3}$  from GEOS-Chem, as well as applying latitudinal gradients to  $P$  and  $I$  based on their values at the three sonde release sites, and adding longitudinal variability based on seasonal stratospheric wind jet streams (Baray et al. 2012; Škerlak et al. 2015). An improved estimate of event lifetime and parameterisation of how many events may be occurring simultaneously could also be addressed, however this is beyond the scope of this work and in any case would not address all the limitations of the estimate provided below.

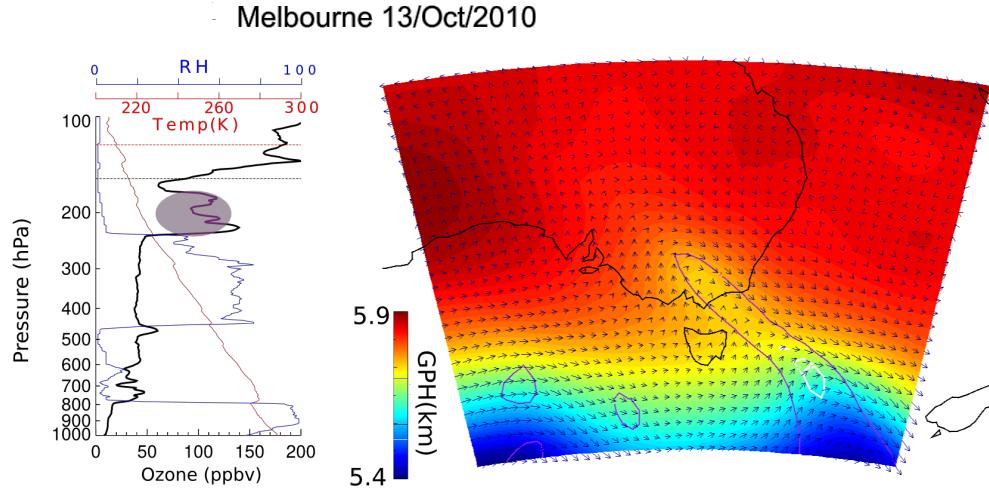


FIGURE S3: Same as Fig. S2 but for 13 January 2010. Also shown in this figure is the 2 PVU contour (white), often used to determine dynamical tropopause height.

### S3.2 Results

Fig. S5 (lower panel) shows the results of the calculation when we choose two days for our flux estimation, with the range shown in representing the values calculated if we assume events last one day (upper bound of estimated flux) or one week (lower bound of estimated flux). Previous studies have found STT ozone fluxes in the SH extratropics are largest from autumn or winter to early spring (Olsen 2003; Škerlak et al. 2015; Liu et al. 2016a). Although these are based on dominating STT systems further north than the area we examine here, see the main text for more details. During the SH winter, we find the highest tropospheric  $\Omega_{O_3}$  but a relatively low STT flux due to reduced event frequency. Our results suggest instead that the ozone flux associated with STT events (at least those due to tropopause folds) is largest in austral summer (December–March), primarily due to an increased frequency of STT detections during these months. It is possible that our estimated event frequencies are too low in late winter–early spring as some legitimate STT events may have been excluded due to coincident smoke plumes.

Summing the monthly estimated fluxes shown in Fig. S5 over the year, we find from this estimate that STT events may be responsible for  $\sim 7.5 \times 10^{16}$  molecules  $\text{cm}^{-2} \text{yr}^{-1}$  of the tropospheric ozone over the Southern Ocean, equivalent to  $75 \text{ Tg yr}^{-1}$ .  $2-16 \times 10^{16}$  molecules  $\text{cm}^{-2}$  of stratospheric based ozone is estimated over the southern ocean throughout the year. Our estimate is hard to directly compare to prior work that suggests global gross STT fluxes of  $550 \text{ Tg yr}^{-1}$  (Stevenson et al. 2006) and net downward STT fluxes of  $75 \text{ Tg yr}^{-1}$  (Sprenger, Croci Maspoli, and Wernli 2003). This is due to the high uncertainties involved in calculation as well as the specific regions which have few other measurements available.

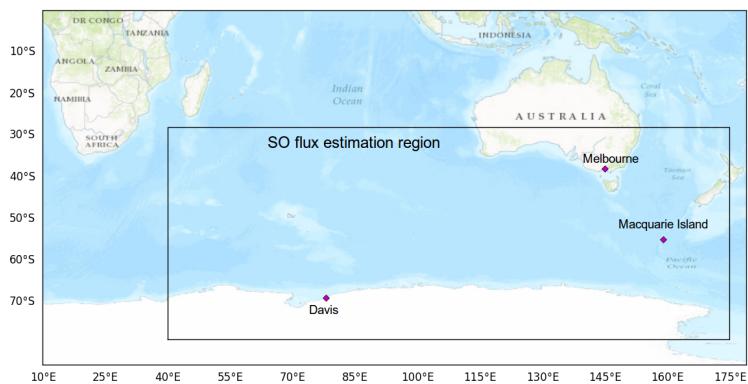


FIGURE S4: Region used for large SO estimation of STT flux

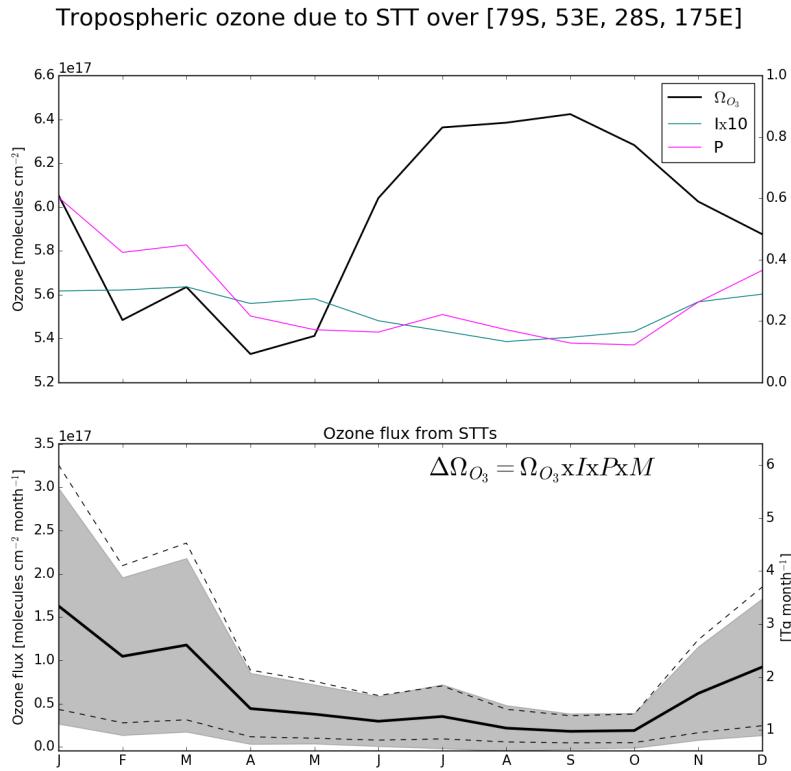


FIGURE S5: (Top) The three quantities used to calculate the total Southern Ocean ozone flux from STT events. The tropospheric ozone column  $\Omega_{O_3}$  (black, left axis) is from GEOS-Chem, while the STT probability  $P$  (magenta, right axis) and impact  $I$  (teal, right axis) are from the ozonesonde measurements. The STT impact is multiplied by 10 to better show the seasonality. (Bottom) Estimated contribution of STT to tropospheric ozone columns over the Southern Ocean. The shaded area shows the uncertainty as calculated in Sect. 6 of the parent document, with the -dashed lines showing the range of values when assuming events last one day (upper dashed line) up to one week (lower dashed line).

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