

SENSITIVITY OF BLACK CARBON AGING TO MODELING ASSUMPTION IN  
CAM-CHEM

BY

YINRUI LI

THESIS

Submitted in partial fulfillment of the requirements  
for the degree of Master of Science in Atmospheric Sciences  
in the Graduate College of the  
University of Illinois at Urbana-Champaign, 2017

Urbana, Illinois

Master's Committee:

Associate Professor Nicole Riemer  
Professor Donald Wuebbles

# Abstract

Black carbon (BC) aerosol strongly absorbs visible light and therefore has a warming impact on climate. Quantifying this impact requires us to develop faithful model representations of its climate-relevant properties, such as CCN activity and optical properties. One key process that needs to be captured is the BC aging process, that is the conversion of fresh, hydrophobic black carbon into aged, hydrophilic black carbon, which directly contributes to CCN activation and wet removal and impacts black carbons optical properties. In current models, the BC aging timescale is either assumed to be a fixed value (1-2 days) or is determined with mechanistic transfer rates based on ad hoc aging aging criteria. Both approaches are very sensitive to the choices of assumed parameters.

The goal of this study is to explore the sensitivity of the simulated BC burden and BC radiative forcing to the aging criterion used in CAMChem, and to compare BC aging rates in CAMChem to an aging parameterization based on more detailed particle-resolved simulations with PartMC-MOSAIC. We carried out a 1-year simulation with the global CAMChem model, where a 4-mode modal aerosol model is used. BC aerosol is transferred from a fresh, hydrophobic mode (primary carbon mode) to an aged, hydrophilic mode (accumulation mode) after condensing a certain amount of secondary materials or through coagulation. Our results show that the simulated BC burden is most sensitive to the choices of the aging criterion in the high-latitude regions, with maximum differences in the annual averaged BC mixing ratio of 16% near the surface. These differences can be higher in the monthly averaged BC mixing ratio (e.g., we observe 65% relative difference at an elevation of 10km in the Arctic for March). We also noted that SP2 instrument can capture most of BC when

it is close to the source regions (internally mixed), and in the Arctic (externally mixed). The aging timescales in the CAMChem model range from less than one hour (South America) to several days (over the ocean) and these values are broadly consistent with the aging timescales from the PartMC-MOSAIC parameterization.

*To my dearest parents for their love and support.*

# Acknowledgments

I would like to express my deepest gratitude to my advisor Prof. Nicole Riemer, who give me strong support and plentiful ideas in this project. I couldn't have accomplish this work without her encouragement and insights. I also want to thank Professor Donald Wuebbles for co-advising this project, and for providing his continuous support.

I'm thankful to the research scientists at the Los Alamos National Laboratory, Manvendra Krishna Dubey and Allison C Aiken, for collaborating on this project and sharing valuable ideas with us. Also I'm grateful to Laura Fierce, whose previous research findings shed an interesting light on our quantification of the carbonaceous aerosol aging timescales, for her enlightening guidance, and Dr. Steven Ghan at the Pacific Northwest National Laboratory, for inviting me to their Lab and helping me through the hard work of model coding.

This work was supported by the Center for Space and Earth Sciences, Los Alamos National Laboratory, Laboratory Directed Research and Development Program. So I would also like to thank them for financially supporting our project, and giving me the opportunity to work on this project.

Lastly, I would like to thank my parents for their love and accompany all the time.

# Table of Contents

<b>List of Figures . . . . .</b>	<b>ix</b>
<b>List of Abbreviations . . . . .</b>	<b>xii</b>
<b>Chapter 1 Introduction . . . . .</b>	<b>1</b>
<b>Chapter 2 Background and Motivation . . . . .</b>	<b>4</b>
2.1 Atmospheric Aerosol and its Particle Size Distribution . . . . .	4
2.2 Composition of Atmospheric Aerosol and its Mixing State . . . . .	6
2.2.1 Chemical Composition of Atmospheric Aerosol . . . . .	6
2.2.2 Aerosol Mixing States . . . . .	7
2.3 Measurements of Carbonaceous Aerosols . . . . .	11
2.4 Model Simulations on Aerosol Evolution . . . . .	13
2.4.1 Aerosol Dynamic Modeling . . . . .	13
2.4.2 Modal Aerosol Model (MAM) in CAM-chem . . . . .	18
2.4.3 Particle Resolved Aerosol Modeling . . . . .	18
<b>Chapter 3 Methodology . . . . .</b>	<b>20</b>
3.1 CAM-chem Model Configurations and MAM4 . . . . .	20
3.2 8-monolayer of Sulfate Condensation Criterion . . . . .	22
3.3 CAM-chem Aging Timescale . . . . .	26
3.4 PartMC-MOSAIC Aging Timescale . . . . .	27
<b>Chapter 4 Results . . . . .</b>	<b>30</b>
4.1 BC Emissions . . . . .	30
4.2 Sensitivity of BC Burden and Climate Effect to the Number of Monolayers .	31
4.2.1 Horizontal Profiles of BC burdens . . . . .	32
4.2.2 Vertical Profiles of BC burdens . . . . .	33
4.2.3 BC Deposition Flux . . . . .	35
4.2.4 Monolayer Effect on Radiative Forcing . . . . .	37
4.3 Mixing State of Species in MAM4 . . . . .	38
4.3.1 Volume Fraction of Species in the Primary Carbon Mode and Accumulation Mode . . . . .	38
4.3.2 BC Mixing States Sensitivity to the Aging Criterion . . . . .	41
4.4 Mass Fraction and Mixing States Observed by SP2 measurement . . . . .	43

4.4.1	BC Core Diameter . . . . .	44
4.4.2	Volume Fraction Within the Size Range of SP2 Measurement . . . . .	45
4.5	BC Mass Fraction Activated by Coating Material . . . . .	49
4.6	Process Analysis of BC Aging . . . . .	50
4.6.1	BC Condensation Growth Rate . . . . .	52
4.6.2	CAM-chem v.s. PartMC–MOSAIC Aging Timescales . . . . .	52
<b>Chapter 5</b>	<b>Summary and Conclusion . . . . .</b>	<b>63</b>
<b>Chapter 6</b>	<b>Appendix . . . . .</b>	<b>66</b>
6.1	Configuration of CAM-chem Model . . . . .	66
<b>References . . . . .</b>		<b>69</b>

# List of Figures

2.1	Overall life cycle of atmospheric aerosols . . . . .	5
2.2	Atmospheric aerosols and their size distributions . . . . .	6
2.3	Chemical species mixing and compositions of aerosol particles, taken from the . (Li et al., 2011b) . . . . .	8
2.4	Location of the AMS datasets analyzed in Zhang et al. (2015). Pie charts show the average mass concentration and chemical composition: organics (green), sulfate (red), nitrate (blue), ammonium (orange), and chloride (purple) . . . . .	9
2.5	Mixing states diagram to illustrate the relationship between per-particle diversity $D_\alpha$ , bulk diversity $D_\beta$ , and mixing state index $\chi$ (Riemer and West (2013)) . . . . .	10
2.6	Diversity and mixing state evolution for BC-containing particles in the urban plume case. (a) Distribution of per-particle diversity $D_i$ as a function of time. (b) Time series of average per-particle diversity $D_\alpha$ , bulk diversity $DD_\gamma$ , and mixing state index $\chi$ (Riemer and West (2013)) . . . . .	11
2.7	Schematic of the Single Particle Soot Photometer (SP2) (Schwarz et al. (2010)).	13
2.8	The average detection efficiencies in each rBC mass-bin for each laser intensity shown in the legend. Whiskers represent the standard deviation of the values in each mass bin. The top axis shows volume-equivalent diameter (assuming 2 g/cc void-free density for rBC) that corresponds to denuded rBC-core mass on the bottom horizontal scale (Schwarz et al. (2010)) . . . . .	14
2.9	Aerosol Size Distribution. ( <a href="http://www.ems.psu.edu/~lno/Meteo437">www.ems.psu.edu/~lno/Meteo437</a> . . . . .	15
3.1	Schematic of aerosol modes and associated tracers in MAM4 . . . . .	22
3.2	Schematic of 8-monolayer of sulfate criterion . . . . .	23
3.3	Raw data that is used to construct the regression function (grey dots) and the resulting regression function (black line) for simulations including only coagulation, without condensation (Fierce et al. (2016)) . . . . .	29
3.4	Estimated e-folding time for the error in CCN activity from the internal mixture approximation. (Fierce et al. (2016)) . . . . .	29
4.1	Monthly mean forest fire and grass fire fluxes of BC for March (top panels) and September (bottom panels), at heights of 0–126 m (left panels) and 278–454 m (right panels) . . . . .	31
4.2	Annual mean anthropogenic emissions of BC emitted at the surface for the year 2010 for 60–90 °N (left) and 70–90 °N (right) . . . . .	32

4.3	Annual BC mass mixing ratio (top) and relative differences denoted by (L1 - L8)/L8 (bottom) . . . . .	33
4.4	Horizontal plots of BC concentrations near the surface for different seasons (March and September) . . . . .	34
4.5	Horizontal map of six regions . . . . .	35
4.6	Vertical profiles of mean burdens for six regions corresponding to Figure 4.5 in March, 2010 . . . . .	36
4.7	Vertical profiles of mean BC burdens for six regions corresponding to Figure 4.5 in September, 2010 . . . . .	37
4.8	Horizontal plots of regional BC concentrations at 859 hPa covering region 1 in case L1 (top) and L8 (middle), and the relative differences (bottom) denoted by (L1 - L8)/L8 . . . . .	38
4.9	Horizontal plots of the relative differences in BC wet deposition flux with different aging criterion (L1 v.s. L8) for March (left) and September (right). . . . .	39
4.10	Horizontal plots of the relative differences in BC dry deposition flux with different aging criterion (L1 v.s. L8) for March (left) and September (right) . . . . .	39
4.11	Horizontal plots of the direct radiative forcing of BC aerosols with different aging criterion (L1 v.s. L8) near the surface (992hPa) . . . . .	40
4.12	Polar plots of the direct radiative forcing of BC aerosols with different aging criterion (L1 v.s. L8) near the surface (992hPa) . . . . .	41
4.13	Annual volume fraction of sulfate, secondary organic aerosol (SOA), black carbon (BC) and organic carbon (OC) near the surface (992hPa) in the accumulation mode . . . . .	42
4.14	Annual volume fraction of dust and sea salt (SST) near the surface (992hPa) in the accumulation mode . . . . .	43
4.15	Annual volume fraction of black carbon (BC) and organic carbon (OC) near the surface (992hPa) in the primary carbon mode . . . . .	43
4.16	Fraction of BC mixing ratio in the accumulation mode to total BC mixing ratio (in %) for L1 case (top panels) and L8 case (bottom panels) in March (left panels) and September (right panels) . . . . .	44
4.17	BC mass fraction within the SP2 size range for primary carbon mode $F_{pc}(d_1, d_2)$ . . . . .	47
4.18	BC mass fraction within the SP2 size range for accumulation mode $F_{accu}(d_1, d_2)$ . . . . .	48
4.19	The ratios of BC mass in the SP2 size range in each mode to the total BC mass in the SP2 size range, for primary carbon mode $f_{pc}(d_1, d_2)$ and for accumulation mode $f_{accu}(d_1, d_2)$ . . . . .	49
4.20	BC mass mixing ratio ( $M_{pc}$ and $M_{accu}$ ) in primary carbon mode (top) and in accumulation mode (bottom), for surface layer, March and September . . . . .	50
4.21	BC mass fraction between 90 and 400 nm ( $F_{pc}$ and $F_{accu}$ ) in the primary carbon mode (top) and accumulation mode (bottom), for surface layer, in March and September . . . . .	54
4.22	Geometric mean diameter of BC core ( $\mu\text{m}$ ) in primary carbon mode (top) and accumulation mode (bottom), for surface layer, in March and September . . . . .	55

4.23 Ratio of BC mixing ratio within SP2 size range to total BC mixing ratio within SP2 size range in primary carbon mode $f_{pc}$ (top) and in accumulation mode $f_{accu}$ (bottom), for surface layer, in March and September . . . . .	56
4.24 Fraction of BC mass that is activated at 0.3% (top) and 0.6% (bottom) supersaturation ratio . . . . .	57
4.25 Volume condensation rate ( $I_{cond}$ ) of sulfate and SOA . . . . .	58
4.26 Annual BC aging timescales (in hours) near the surface, extracted from the CAM-chem model (left) and computed based on PartMC-MOSAIC parameterization (right). The condensation, coagulation and overall timescales are displayed in the upper panels, middle panels and bottom panels respectively . . . . .	59
4.27 Annual BC aging timescales (in hours) at 857 hPa, extracted from the CAM-chem model (left) and computed based on PartMC-MOSAIC parameterization (right). The condensation, coagulation and overall timescales are displayed in the upper panels, middle panels and bottom panels respectively . . . . .	60
4.28 Scatter plots of CAM-chem v.s. PartMC–MOSAIC aging timescales (in hour) near the surface in 5 sensitivity cases. L1–L8 represents the number of monolayers (1 to 8) for the condensation criterion . . . . .	61
4.29 Scatter plots of CAM-chem v.s. PartMC–MOSAIC aging timescales (in hour) at 857 hPa in 5 sensitivity cases. L1–L8 represents the number of monolayers (1 to 8) for the condensation criterion . . . . .	62

# List of Abbreviations

BC	Black Carbon.
EC	Elemental Carbon.
OC	Organic Carbon.
PM	Particulate Matter.
VOC	Volatile Organic Components.
SOA	Secondary Organic Aerosol.
CCN	Cloud Condensation Nuclei.
IN	Ice Nuclei.
TEM	Transmission Electron Microscopy.
GDE	General Dynamic Equation.
MDE	Moment Dynamic Equation.
MAM <sub>n</sub>	n-mode Modal Aerosol Model.
SP2	Single Particle Soot Photometer.

# Chapter 1

## Introduction

Black carbon (BC) aerosol is a product of incomplete combustion of fossil fuel, biofuel and biomass burning (e.g., Bond et al. (2004); Forsström et al. (2013)). It strongly absorbs visible light and has been ranked as the second most important individual absorbing agent after CO<sub>2</sub>, with a climate forcing of +1.1Wm<sup>-2</sup> (Bond et al. (2013)). BC aerosols can impact the global atmospheric radiative budget by absorbing or scattering incoming solar radiation, which is called the direct effect. ~~When solar radiation is absorbed by aerosols, there is often a warming effect, whereas when the radiation is scattered back, there is often a net cooling effect.~~ BC aerosols can also impact the climate by acting as cloud condensation nuclei or ice nuclei and affect cloud microphysical properties. This effect is referred to as the indirect effect. Furthermore, it can act as impurities in snow and ice after being transported to the polar regions and depositing onto their surfaces, ~~resulting in a positive feedback on the surface albedo~~ (Zuberi et al. (2005); Flanner et al. (2007)). BC aerosols also have adverse impacts on air quality and human health (Highwood and Kinnersley (2006)).

Quantifying these impacts requires us to develop faithful model representations of the BC burden and its climate-relevant properties such as cloud condensation nuclei (CCN) activities and optical properties. Currently, however, significant discrepancies in model simulated BC remain in most global climate models (GCMs) and large uncertainties of its climate forcing have been shown. For example, previous studies have implied a general overestimation of BC in the mid-upper troposphere in the mid-latitudes, and an underestimation of BC in the lower and middle troposphere at high latitude (Koch et al. (2009); Schwarz et al. (2010); Fan et al. (2012)). The simulated BC aerosol absorption optical depth tends to be biased low

compared to satellite observations (e.g., Koch et al. (2009)). These model failures can result from the complex BC aerosol processes and properties that has not been well captured, such as its emissions, coagulation, condensation, dry deposition, wet scavenging, etc. (Hakami et al. (2005); Koch et al. (2009); Shindell et al. (2012)). These processes in turn control the evolution of aerosol burden, size distribution, mixing states (the level to which particles are internally or externally mixed with other species or in between), and consequently its climate forcing (e.g., Schulz et al. (2006)).

Among them, one key process that contributes to the uncertainties and thus needs to be captured is the BC aging process, the conversion of freshly emitted, hydrophobic BC to aged, hydrophilic BC through coating with sulfate and organics or coagulation (Langner (1992); Parungo et al. (1994); Liousse et al. (1996)). This process directly contributes to the CCN activation and wet removal, and also impacts black carbon's optical properties by evolving the composition and mixing states of aerosols. Therefore, it plays a significant role in simulating the lifetime of BC, and hence its transport, distributions and climate effects (Croft et al. (2005); Riemer et al. (2004)). In addition, previous studies have found that the parameterizations of the aging process can significantly affect model results (Liu et al. (2010)).

Albeit its importance, the treatment of BC aging, however, is usually very simplified in global scale models by using fixed timescales or parameterized aging rates for the sake of computational limits. The most simplified bulk scheme assumes fully externally mixed populations and often use a fixed aging timescale (on the order of 1-2 days) for conversion of hydrophobic BC to hydrophilic BC (Cooke and Wilson, 1996). There are also more advanced and complicated schemes such as the modal aerosol model or sectional models, computing mechanic transfer rates by assuming aerosol size distributions and mixing levels (Bauer et al. (2013), Huang et al. (2013)).

The representation of BC aging in the Community Atmosphere Model with Chemistry (CAM-Chem), an atmospheric component of the Community Earth System Model (CESM),

uses the latter scheme. It applies a 4-mode version of the ~~modal aerosol model~~ (MAM4), where BC is emitted to the primary carbon mode, and then is aged and transferred to the accumulation mode by condensation of sulfate, ammonia and SOA and by coagulation (Liu et al. (2012); Lamarque et al. (2012)). In MAM4, a criterion of 8 monolayers of sulfate is used to compute the aerosol transfer rate from primary carbon mode to accumulation mode. It assumes that BC particle is aged after condensing an equivalent of 8 monolayers of sulfate. However, previous studies have shown that considerable sensitivities exist regarding the choices of the number of monolayers and other parameters, and significant model biases in BC burden have been found compared to HIPPO observations (Liu et al. (2010)).

Furthermore, Fierce et al. (2016) has derived a parameterization that characterizes the aging rates of BC aerosols through gas condensation and particle coagulation from detailed simulations on the particle scale, based on the particle-resolved PartMC-MOSAIC model (Particle Monte Carlo Model for Simulating Aerosol Interactions and Chemistry) (Fierce et al. (2016)). PartMC-MOSAIC is a complex aerosol model that provides detailed information on aging processes at the micro-scale, by tracing the size, composition and mixing states of particles (Riemer (2010)). The aging parameterization put forward by Fierce et al. (2016) can be used to evaluate and to improve the treatment of BC aging in the global-scale models.

In this work, our aim is to assess the representation of BC ~~aerosols~~ in MAM4 of CAM-Chem model. We conduct ~~several sensitivity runs of BC mixing ratio, mixing states and direct radiative forcing to its condensation criterion~~, in order to investigate the extent to which ~~those quantities~~ are sensitive to the choices of aging parameters. We also exploit the PartMC-MOSAIC parameterization as the reference to evaluate the performance of MAM4 aging scheme.

# Chapter 2

## Background and Motivation

In this chapter, starting with an introduction to atmospheric aerosols, and more specifically the BC-containing aerosols, we give a broad view of their size distribution, chemical compositions and mixing state, in preparation for the further discussion of their measurements and representation in numerical models. We then explained the mixing state impact on climate and the benefit and limit of SP2 measurements. Model simulation of aerosols is discussed at the end of this chapter, including the general aerosol dynamic modeling, CAM-chem modal aerosol modeling (MAM) and Particle Resolved Aerosol Modeling.

### 2.1 Atmospheric Aerosol and its Particle Size Distribution

Atmospheric aerosol is a multiphase system containing gases with suspended particles in the air. They can come from either natural sources or anthropogenic sources, such as dust and sea salt that arises without human intervention, pollen and plant debris that originate from life processes, and vehicle and industrial combustions that come from human activities. Figure 2.1 illustrates the aerosol life cycle in the atmosphere, including their sources, physical and chemical processes during their formation and transport, and their deposition.

As a result, aerosols can change the composition of the atmosphere and cause some environmental and climate effects. For example, carbonaceous aerosol has been associated with natural and anthropogenic air pollution, reduced visibility, climate change, and ad-

verse health effects (Mauderly and Chow (2008)). One of the most important parameters that affect the transport scale and the environmental and climate properties of atmospheric aerosols is the size of their particles (Pacyna (1995)). Figure 2.2 shows the size distributions of the equivalent aerodynamic diameter of aerosols (the diameter of a spherical particle with the same value of some physical property as the irregular particle), and their categorization into modes with respect to their sizes and origins. Generally, nucleation mode consists of particles smaller than  $10^{-2}\mu\text{m}$ , formed by condensation of supersaturated vapors or through gas-to-particle conversion. They grow in size by condensation of vapor and coagulation with other particles, and can form Aitken mode ( $10^{-2}\mu\text{m} - 10^{-1}\mu\text{m}$ ) and accumulation mode ( $10^{-1}\mu\text{m} - 1\mu\text{m}$ ). Particles above the diameter size of  $1\mu\text{m}$  form the coarse mode. They are produced by mechanical processes such as the fragmentation of solids and atomization of liquids (Pacyna (1995); Kulmala et al. (2004)). The nucleation mode and Aitken mode account for most of the number of the aerosol population, whereas the accumulation and coarse mode account for most of the mass.

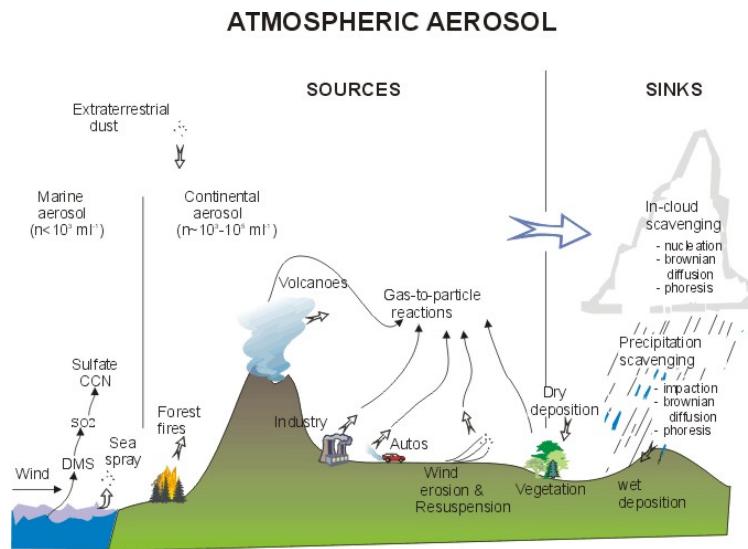


Figure 2.1: Overall life cycle of atmospheric aerosols.  
<http://www.ems.psu.edu/lno/Meteo437/Aerosol.jpg>

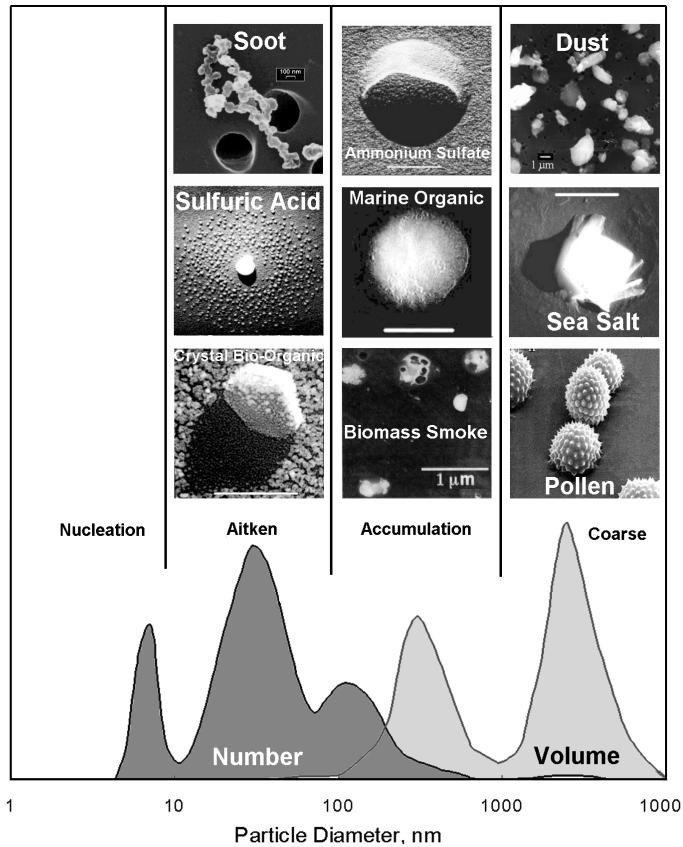


Figure 2.2: Atmospheric aerosols and their size distributions.(<http://capita.wustl.edu>)

## 2.2 Composition of Atmospheric Aerosol and its Mixing State

### 2.2.1 Chemical Composition of Atmospheric Aerosol

Atmospheric aerosols can be formed either by direct emissions from sources such as incomplete combustion and wind-driven suspension of solids (primary particle), or by gas-to-particle conversion (secondary particle). Sulfate, nitrate, ammonium, chloride, sea salt, mineral dust, organic components and black or elemental carbon are the predominant components of the air particulate matters (PM) (Pöschl (2005)).

Organic species represent one major fraction (20–90 %) of the aerosol mass. The primary

organic aerosols are emitted directly as liquid or solid particles or as semivolatile vapors, and the secondary organic aerosols are formed by gas-to-particle conversion of volatile organic components (VOCs). The main sources of organic species include natural or anthropogenic biomass burning, fossil-fuel combustion and biological materials (plants, pollen, animal debris)(Pöschl (2005)). Another main components of air aerosol mass are carbonaceous species that consist of black carbon (BC) or elemental carbon (EC) and organic carbon (OC). Freshly emitted BC particles are mostly hydrophobic, and the particles tend to grow hygroscopic via condensation of secondary organic and inorganic species and coagulation with soluble aerosols (Riemer et al. (2010)). The composition of carbonaceous aerosols is dependent on the fuel type and

Figure 2.3 shows an observation of the chemical species compositions of single aerosol particles, measured by transmission electron microscopy (TEM). It is obvious that most aerosols are not pure, and can be mixed with other species. In addition, the composition can be highly variable spatially. Figure 2.4 shows the average mass concentration and chemical composition of the non-refractory submicron particles based on the AMS datasets.

### 2.2.2 Aerosol Mixing States

We have mentioned in the previous section that most aerosols are not pure, and can be mixed with other species. The degree to which particles are internally or externally mixed with other species or in between are called 'mixing state'.

#### A Quantitative Metric

Riemer and West (2013) has developed the first quantitative metric to represent the mixing state of aerosol population using a mixing ratio index  $\chi$  defined as the ratio of per particle diversity  $D_\alpha$  to the bulk population diversity  $D_\gamma$ . It ranges from  $\chi = 0$  when all particles are fully externally mixed to  $\chi = 1$  when all particles are fully internally mixed. Figure 2.5 illustrates the relationship between per-particle diversity  $D_\alpha$ , bulk diversity  $D_\beta$ , and mixing

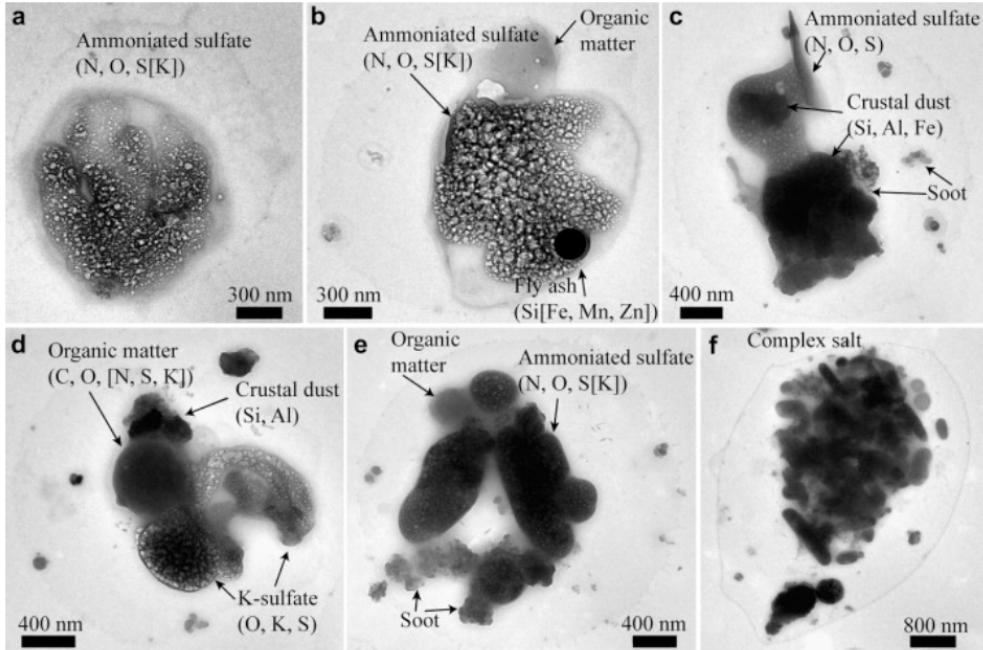


Figure 2.3: Chemical species mixing and compositions of aerosol particles, taken from the . (Li et al., 2011b)

state index  $\chi$ .

The initial mixing state can be further modified in the atmosphere as particles evolve. Taking carbonaceous aerosols for example, freshly emitted BC aerosols are mostly hydrophobic and externally mixed. During their transport, they tend to become more hydrophilic and internally mixed as a result of the aging process –the condensation of secondary organic and inorganic species and coagulation with soluble aerosols. Figure 2.6 shows the evolution of the carbonaceous aerosol mixing states with time. As a result, the accuracy of BC mixing state and related climate properties greatly depend on the representation of the aging process in models.

### Mixing State Impact on Climate

Current representation of aerosol and its impact on climate in global models still has large uncertainties, mostly due to the difficulties to capture the microphysical and chemical pro-

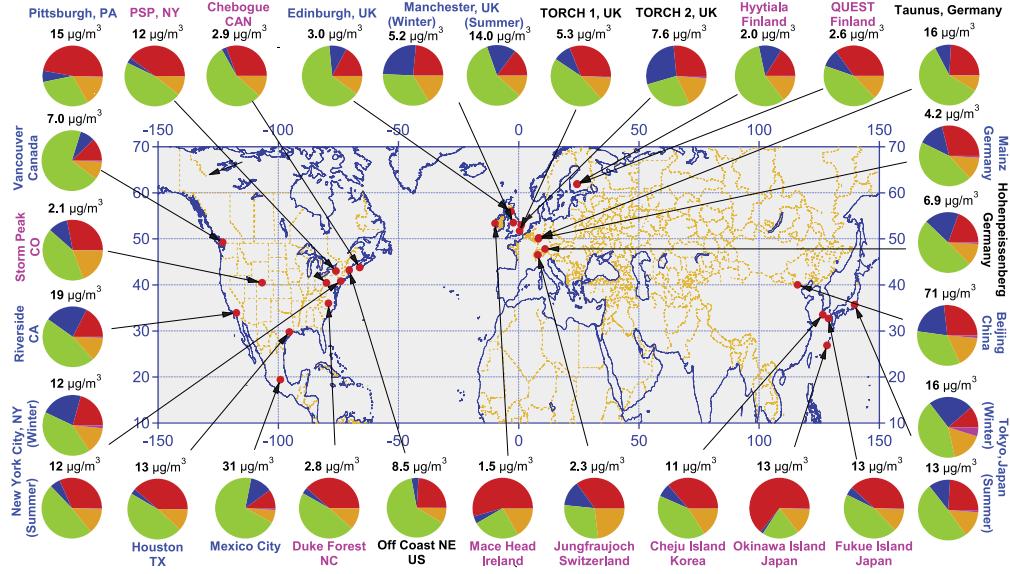


Figure 2.4: Location of the AMS datasets analyzed in Zhang et al. (2015). Pie charts show the average mass concentration and chemical composition: organics (green), sulfate (red), nitrate (blue), ammonium (orange), and chloride (purple).

cesses that drive the evolution of aerosol particles in the atmosphere. Several studies have implied that the mixing states affect the climate-related aerosol properties such as optical properties or cloud condensation nuclei activity (Jacobson (2001); Zaveri et al. (2010); Koch et al. (2009)). In addition, Reddington et al. (2013) has found that the model-observation biases in BC properties are much greater than for the overall particle distribution, indicating that the model discrepancies can be largely due to the assumptions about the size and mixing state of the emitted carbonaceous particles.

Efforts to improve the representation of aerosol mixing states in both regional and global models have also been made (e.g., Jacobson (2001); Riemer and West (2013)). Riemer and West (2013) has pointed out that the mixing state of carbonaceous particles depends strongly on the fuel and combustion type. In our study, we computed the average population hygroscopicity  $\kappa$  as an implication to mixing states for each mode of the 4-mode modal aerosol model (MAM4) in CAM-chem.

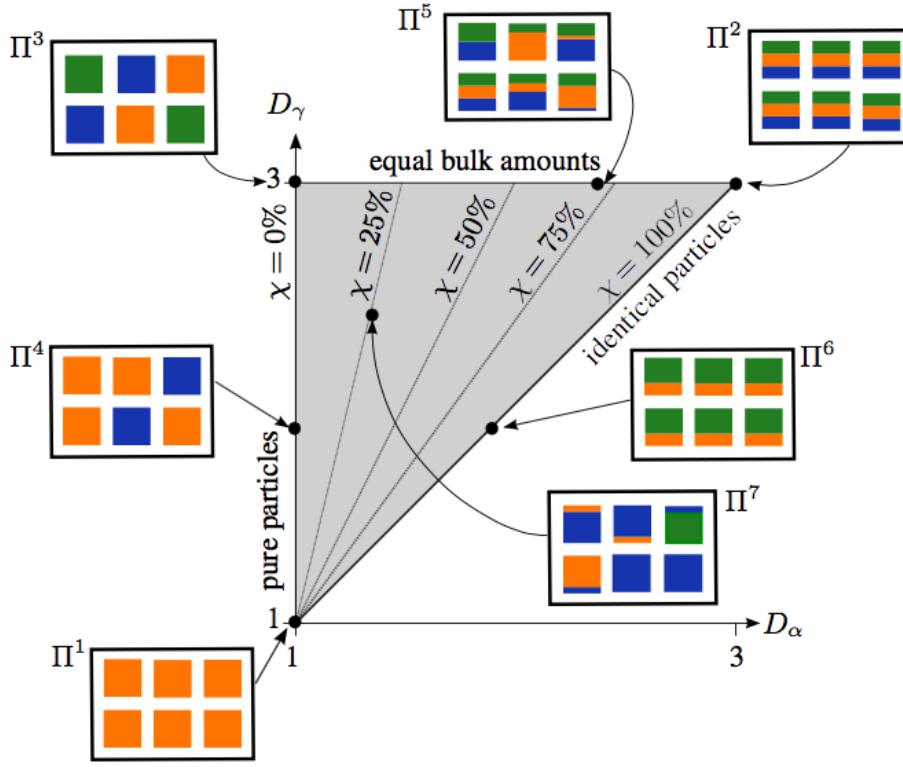


Figure 2.5: Mixing states diagram to illustrate the relationship between per-particle diversity  $D_\alpha$ , bulk diversity  $D_\beta$ , and mixing state index  $\chi$  (Riemer and West (2013)).

### BC-containing Aerosols

BC-containing aerosols can affect climate in multiple ways. They have both direct and indirect effects as atmospheric aerosols, and can reduce the surface albedo after depositing onto snow or ice. The BC component of its aerosols strongly absorbs solar radiation at visible wavelength, however, it can also be mixed with hydrophilic materials such as [ammonia](#), sulfate or secondary organic aerosols (SOA) and act as CCN or IN. The mixing state of atmospheric BC particles with those hydrophilic aerosol components can affect both their CCN/IN activities and radiative properties (e.g., Cheng et al. (2006)). Previous study has found that the light absorption of BC particles can be enhanced by 50 to 60 % after adding non-BC materials, and the enhancement is determined by the particles' mass ratio of non-

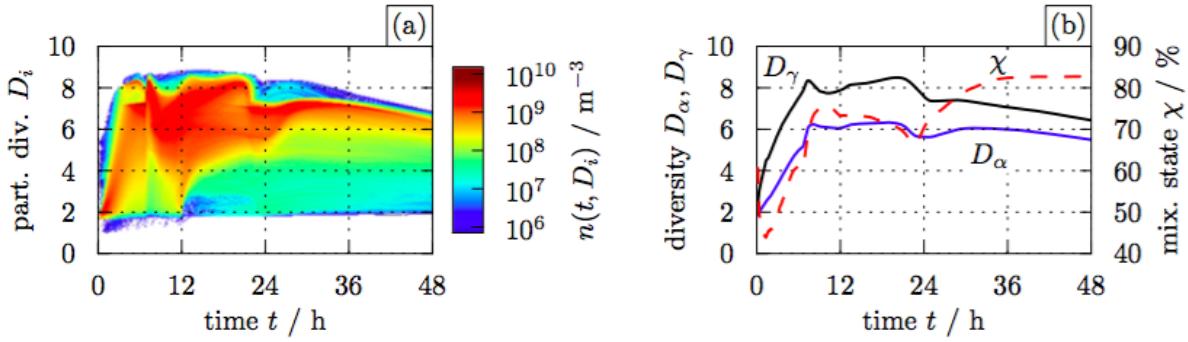


Figure 2.6: Diversity and mixing state evolution for BC-containing particles in the urban plume case. (a) Distribution of per-particle diversity  $D_i$  as a function of time. (b) Time series of average per-particle diversity  $D_\alpha$ , bulk diversity  $D_\gamma$ , and mixing state index  $\chi$  (Riemer and West (2013))

black carbon (Liu et al. (2017)).

In addition, current atmospheric models used to quantify the climate effects of black carbon aerosols are to be improved due to the uncertainties in simulating the complex processes such as BC aerosol formation, aging, cloud interaction, and deposition (e.g., Kipling et al. (2016)). This implies that the model predicted BC aerosol mixing state are generally not in agreement with the observations (e.g., Raatikainen et al. (2015)).

## 2.3 Measurements of Carbonaceous Aerosols

State measurements are required in order to assess the accuracies of model simulated BC aerosol concentrations, mixing states and climate effect, and to improve model parameterizations (Reddington et al. (2013); Reddington et al. (2013)). However, though many measurements of BC aerosols have been launched, few instruments are able to measure their mixing state. In recent years, one advanced observation technique using laser-induced incandescence, called the Single Particle Soot Photometer (SP2), has been developed to measure the mass distribution and mixing state of refractory black carbon (rBC) (Baumgardner et al. (2004)); Schulz et al. (2006)). It is probably one of the most widely used instruments for

BC aerosols.

Single particle soot photometer (SP2) quantifies refractory black carbon (rBC) mass by using a laser to heat rBC to evaporation and measuring their emitted thermal radiation (Figure 2.7). It measures the BC particle cores over a calibrated volume equivalent diameter (VED) range of 55–400 nm. This method is suited for the detection of BC mass in the accumulation mode (Schwarz et al. (2010)), but unlikely to represent the total ambient number and mass concentrations of BC particles. The SP2 number-detection efficiency at sea level pressure is reported to be 100% for BC above 90 nm VED (Schwarz et al. (2010), Figure 2.8), so following Reddington et al. (2013), we regarded 90 nm–400 nm as the efficient diameter range.

In order to compare model simulated BC to observations, it is important to first make sure that most of the modeled BC mass is in the size range of SP2 measurements (e.g., in source regions, freshly emitted BC can be smaller than 100 nm). In our study, we estimated the mass fraction of modeled BC in the size range corresponding to SP2 measurement in preparation for further comparison with observations, especially in the Arctic region.

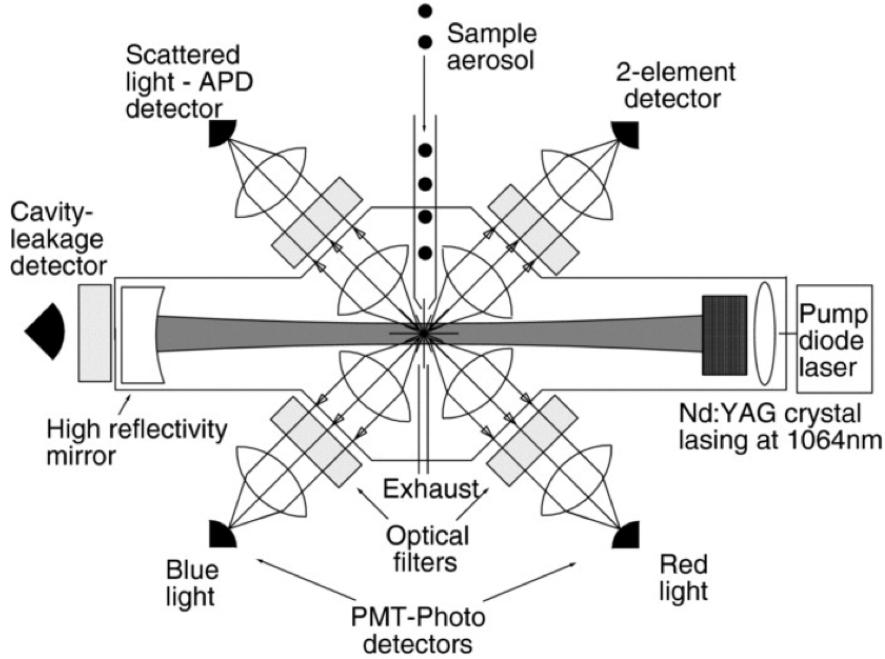


Figure 2.7: Schematic of the Single Particle Soot Photometer (SP2) (Schwarz et al. (2010)).

## 2.4 Model Simulations on Aerosol Evolution

### 2.4.1 Aerosol Dynamic Modeling

During their lifetime in the atmosphere, aerosols undergo complex physical and chemical processes. A numerical aerosol model can serve as useful tool to gain process-level understanding of the aerosol evolution in the atmosphere. Generally, an aerosol model should be able to represent the relevant processes including coagulation, condensation of inorganic and secondary organic species, nucleation, emission and chemical reaction. (Whitby and McMurry (1997)). The approaches to solve those expressions differ by the way that aerosol size distributions are approximated. Some traditional models represent aerosols as a bulk population or as a function of single variable such as their mass. More computationally demanding models include sectional models that sets discrete size bins and typically assumes species to be internally mixed with the pre-existing aerosols in each bins and externally mixed between different bins (Jacobson (2001); Adams et al. (1999)). This approach neglects the aging

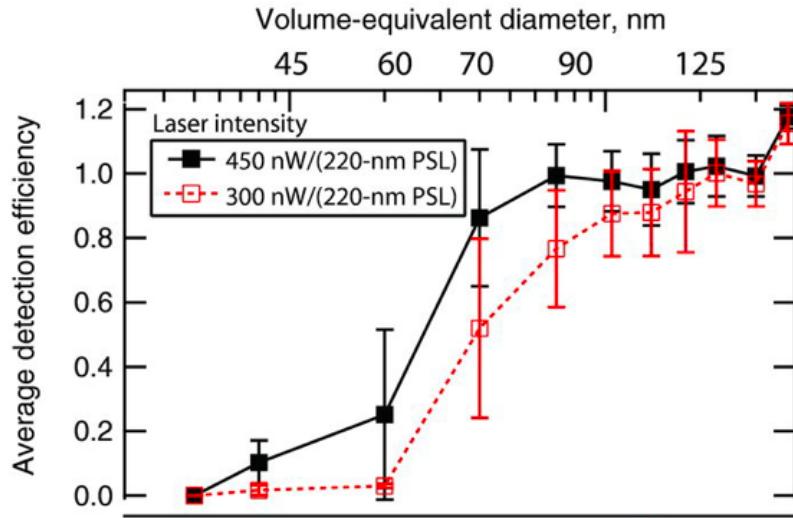


Figure 2.8: The average detection efficiencies in each rBC mass-bin for each laser intensity shown in the legend. Whiskers represent the standard deviation of the values in each mass bin. The top axis shows volume-equivalent diameter (assuming 2 g/cc void-free density for rBC) that corresponds to denuded rBC-core mass on the bottom horizontal scale (Schwarz et al. (2010)).

process by assuming particles to be internally mixed immediately after emission. Another frequently-used representation is called the modal aerosol model (Whitby and McMurry (1997); Binkowski and Shankar (1995)). Aerosols are viewed as distinct populations of particles that is distinguished by their size or chemical composition, and the size distribution of each population is approximated by some analytical distribution function (Binkowski and Shankar (1995)). Each region of the size distribution is called a mode. The size of the particles in each population is typically approximated by a lognormal distribution characterized by the mean diameter  $\mu$  and a standard deviation  $\sigma$ . The accuracy of the model largely depend on how well the size distribution functions of the modes can represent the actual aerosol size distributions.

## Aerosol Size Distribution

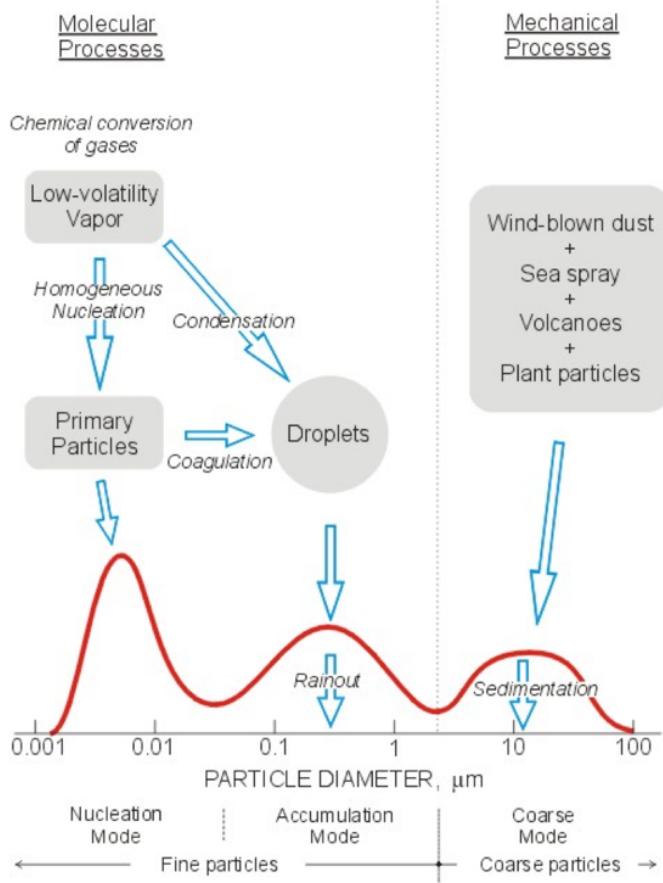


Figure 2.9: Aerosol Size Distribution. ([www.ems.psu.edu/~lno/Meteo437](http://www.ems.psu.edu/~lno/Meteo437))

### Moment and Moment Dynamic Equation

Modal aerosol models typically assume lognormal size distributions for each mode (Equation 2.1) by setting a mean diameter  $D_g$  and a standard deviation  $\sigma_g$ .

$$n(\ln D) = \frac{N}{\sqrt{\pi} \ln \sigma_g} \exp\left[-\frac{1}{2}\left(\frac{\ln(D/D_g)}{\ln \sigma_g}\right)^2\right] \quad (2.1)$$

We can predict how the size distribution evolves if we know how the three parameters,  $D_g$ ,  $N$ , and  $\sigma_g$  evolve in the response to the microphysical and chemical processes such as nucleation, coagulation and condensation. During each time step, the model will update the

geometric mean diameter  $D_g$ , total number concentration  $N$  and standard deviation  $\sigma_g$  for each mode, solved from their aerosol dynamic differential equations.

However, while the differential equations for the total number  $N$  is easy to obtain, it is hard to find the analogous equations for  $D_g$  and  $\sigma_g$ . In order to avoid this problem, some kind of integral form, called moment, is hence introduced to replace the traditional form of differential equations. The mathematical expression of the  $k$ th moment is defined as:

$$M_k = \int_{-\infty}^{+\infty} D^k n(\ln D) d\ln D = N D_g^k \exp\left[\frac{k^2}{2} \ln^2 \sigma_g\right] \quad (2.2)$$

$k = 0$  : total number  $N = M_0$

$k = 2$  : total surface area  $S = \pi M_2$

$k = 3$  : total volume  $V = \frac{\pi}{6} M_3$

—where the total number, surface area and volume of particles in one mode are proportional to the 1st moment, 2nd moment and 3rd moment respectively.

The general dynamic equation (GDE) for aerosols can be found in Whitby and McMurry (1997) (equation 4). Here we only show an analogous form of the general dynamic equation written for the change of moments,  $M_k$ , called the moment dynamic equation (MDE):

$$\begin{aligned}
\frac{\partial M_k(t)}{\partial t} = & \underbrace{\nabla \cdot v M_k}_{\text{advection}} - \underbrace{\nabla \cdot \int_0^\infty D^k \times c(D, t) n(D, t) dD}_{\text{external forces}} \\
& + \underbrace{\nabla \cdot \int_0^\infty D^k \times D(D, t) \nabla n(D, t) dD}_{\text{diffusion}} \\
& + \underbrace{\int_0^\infty \frac{dD^k}{dV(D)} \times \frac{\partial V(D)}{\partial t} \times n(D, t) dD}_{\text{condensation/evaporation}} \\
& + \underbrace{\int_0^\infty D^k \times \dot{n}(D, t) dD}_{\text{nucleation}} \\
& + \underbrace{\frac{1}{2} \int_0^\infty \int_0^\infty (D_1^3 + D_2^3)^{\frac{k}{3}} \times \beta(D_1, D_2) \times n(D_1, t) \times n(D_2, t) dD_1 dD_2}_{\text{coagulation gain}} \\
& - \underbrace{\frac{1}{2} \int_0^\infty \int_0^\infty (D_1^k + D_2^k) \times \beta(D_1, D_2) \times n(D_1, t) \times n(D_2, t) dD_1 dD_2}_{\text{coagulation loss}}
\end{aligned} \tag{2.3}$$

In Equation 2.3,  $c(D, t)$  is the deposition rate,  $D(D, t)$  is the diffusion rate,  $n(D, t)$  is the particle number distribution,  $\beta(D_1, D_2)$  is the coagulation coefficient between particles with geometric size of  $D_1$  and  $D_2$ , and  $V(D)$  is the volume distribution of particles. The coagulation terms only represent intramodal coagulation here. When there are more than 2 modes, the equation will include more coagulation terms in the same form as the fifth and sixth term, in order to represent intermodal gain and loss due to coagulation.

Generally, modal aerosol model will solve the moment differential equations for  $M_0$ ,  $M_3$  and  $M_6$ , and then derive  $N$ ,  $D_g$  and  $\sigma_g$  from the three moments.  $M_6$  is chosen because the coagulation term can be integrated analytically.

## 2.4.2 Modal Aerosol Model (MAM) in CAM-chem

Figure 2.9 shows a typical categorization of aerosols by their size distributions. Particles less than  $0.1 \mu\text{m}$  form the nucleation mode, particles in the size range  $0.1\text{--}2 \mu\text{m}$  form the accumulation mode, and particles larger than  $2 \mu\text{m}$  form the coarse mode. Similarly, in the CAM-chem model, a 3-mode version of modal aerosol model (MAM3) that has only Aitken, accumulation and coarse mode was developed as default, which is efficient for long-term simulations (Liu et al. (2012)). There are also a 4-mode version (MAM4) that has an additional primary carbon mode ~~on top of~~ MAM3 for the treatment of the microphysical aging processes of primary carbonaceous aerosols (Liu et al. (2016)), and a 7-mode version (MAM7) that has Aitken, accumulation, primary carbon, fine dust and fine sea salt, coarse dust and coarse sea salt modes (Liu et al. (2012)). MAM7 has similar treatment of the aging process as in MAM4, but it also separates out sea salt and dust.

We used MAM4 in our study. More description about MAM4 is in section 3.1. MAM4 assumes that the standard deviation  $\sigma_g$  for each mode is fixed, whereas  $N$  and  $D_g$  evolve with time. For each time step the model will solve the dynamic differential equations for  $M_0$  and  $M_3$ , and then derive  $N$  and  $D_g$  from the two moment equations.

## 2.4.3 Particle Resolved Aerosol Modeling

To compensate our study of carbonaceous aerosol aging timescales, we need a more complex and precise model as a reference to the global model simulated results.

A particle resolved Monte Carlo (PartMC) model offers a way to explicitly resolve and tracks the composition of individual aerosol particles in a well-mixed volume (Riemer et al. (2009)). Coagulation, advection and diffusion are simulated stochastically. It is also coupled with the model for simulating aerosol interactions and chemistry (MOSAIC) to simulate the gas-phase chemistry and dynamic gas particle mass transfer (Zaveri et al. (2008)).

PartMC-MOSAIC model predict the evolution of mass and composition of individual

particles, and hence their mixing states and climate related properties (e.g., CCN property, optical depth property) without making simplified assumptions. So the errors of simulated aerosols and their aging processes can be reduced. In this sense, it has the benefit over the traditional sectional or modal models that often make assumptions for aerosol distribution, mixing states and the condensation and coagulation processes.

# Chapter 3

## Methodology

In this chapter, CAM-chem model configuration, an equivalent of 8 monolayer of sulfate condensation criterion for BC aging and its mathematical representation will be discussed. Also, the first-order BC aging timescale extracted from CAM-chem model and the PartMC-MOSAIC parameterization of aging that can be used as a reference for global models will be explained at the end of this chapter.

### 3.1 CAM-chem Model Configurations and MAM4

The CAM-chem model is a global 3-D atmospheric component of the NCAR Community Earth System Model (CESM). It consists of the Community Atmosphere Model (CAM) model and chemical mechanism of a fully implemented model for ozone and related chemical tracers (MOZART-4) which includes 191 chemical tracers and over 400 reactions. CAM-chem model can be run either with interactive meteorology coupled to a free-running ocean with specified sea ice and sea surface temperature (free-running configuration), or with specified meteorology which read in winds, air temperature, surface pressure and heat fluxes (offline configuration) (Lamarque et al. (2012)). It is also coupled to the land model, involving emissions from biogenic sources by the Model of Emissions and Aerosols from Nature (MEGAN).

For our study, we chose the CESM 1-2-2 CAM-chem configured with the offline meteorological data from the Global Earth Observing System (GEOS-5) of the Global Modeling Assimilation Office of NASA, and launched one-year simulation for 2010 with a standard

horizontal resolution of 1.9 latitude by 2.5 longitude and a vertical resolution of 56 layers in order to match the resolution of the input meteorology field. We use the Intergovernmental Panel on Climate Change (IPCC) Fifth Assessment report (AR5) gridded POM and BC emissions for the period 1850-2010 in decadal increments (Lamarque et al. (2010)). This inventory covers a wide range of sources including anthropogenic emissions (without injection heights) originating from domestic, energy, industry, transportation, waste treatment and ship activity sectors, and natural emissions (elevated) from forest fire and grass fire. It assumes that the POM emissions are higher than the OC emissions by a factor of 1.4 (Liu et al. (2012)) and the injection heights for fires are from Dentener et al. (2006). The model treats hydrophobic BC and hydrophilic BC as two separate species, and all the freshly emitted BC falls into the first category.

There are three versions of modal representation of aerosol in CAM-chem. A 3-mode version of modal aerosol model (MAM3) that has only Aitken, accumulation and coarse mode is developed as default, which is efficient for long-term simulations (Liu et al. (2012)). There are also a 4-mode version (MAM4) that has an additional primary carbon mode on top of MAM3 for the treatment of the microphysical aging processes of primary carbonaceous aerosols (Liu et al. (2016)), and a 7-mode version (MAM7) that has Aitken, accumulation, primary carbon, fine dust and fine sea salt, coarse dust and coarse sea salt modes (Liu et al. (2012)). MAM7 has similar treatment of the aging process as in MAM4, but it also separates out sea salt and dust. MAM assumes lognormal size distributions for each mode by setting a fixed standard deviation, and the geometric mean diameter and the total number concentration in each mode can evolve with time. In MAM3, all BC particles are assumed to be aged and put directly into the accumulation mode, where particles are exposed to in-cloud and below-cloud wet deposition. In our study, we applied the newly developed MAM4 model, where the freshly emitted BC/POM particles will come directly into the hydrophobic primary carbon mode. The hygroscopicity of BC is set to be 0, and the hygroscopicity of POM is set to be 0.10, allowing them to experience some in-cloud scavenging in the primary

carbon mode. After the aging process due to condensation of sulfate aerosols, ammonia and some semi-volatile organic aerosols and inter-mode or intra-mode coagulation, the fresh carbonaceous particles can be transferred from the primary carbon mode to the accumulation mode, viewed as being aged. A schematic of aerosol modes and associated tracers in MAM4 is shown in Figure 3.1.

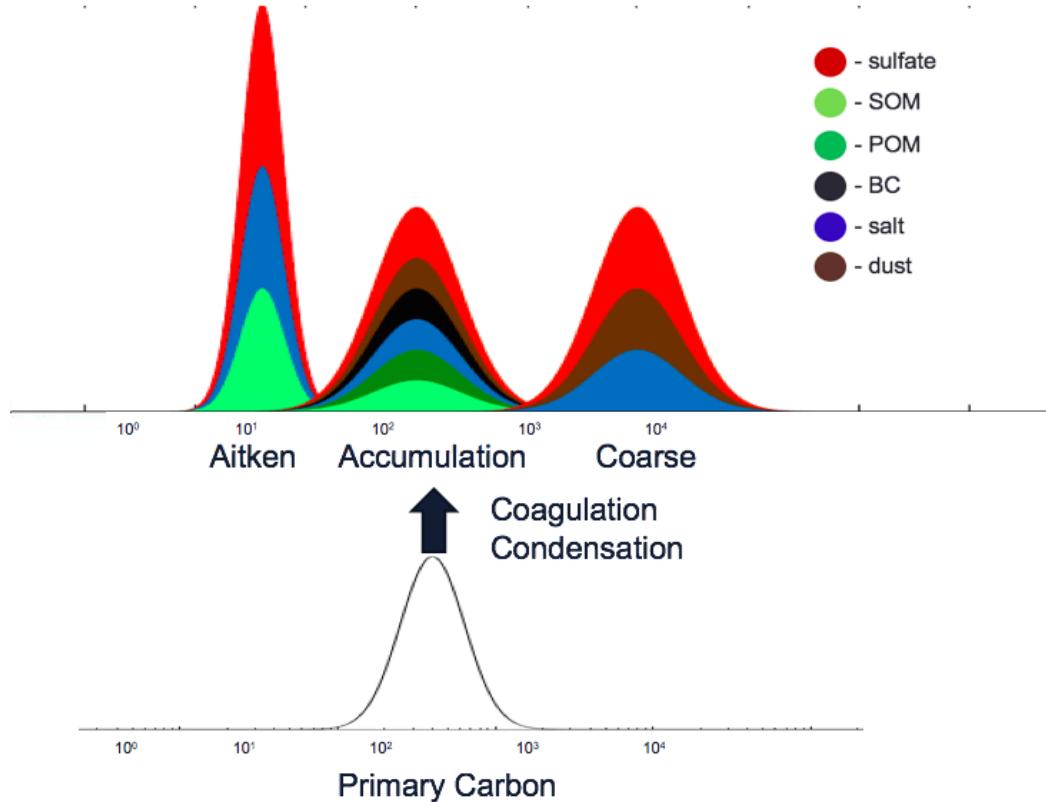


Figure 3.1: Schematic of aerosol modes and associated tracers in MAM4.

## 3.2 8-monolayer of Sulfate Condensation Criterion

In MAM4, condensation of sulfate, ammonia and semi-volatile organics to carbonaceous particles are treated in a dynamic way, where a “monolayer condensation criterion” is applied. It assumes that BC particles become hydrophilic after condensing a equivalent of 8-monolayer

of sulfate onto its core surface, and then this mass will be transferred from the primary carbon mode to the accumulation mode based on the standard mass transfer expressions, regarded as being aged (Liu et al. (2012)). A schematic of the 8-monolayer condensation criterion is shown in Figure 3.2. The depth of one monolayer of sulfate is equal to the molecular diameter of sulfate particle ( $4.76 \times 10^{-10}$  m). It needs to be noted that MAM4 doesn't track individual particles but instead simulate the evolution of the population distribution in each mode. We found that the simulated BC concentration and direct radiative forcing are really sensitive to the choice of the aging criterion.

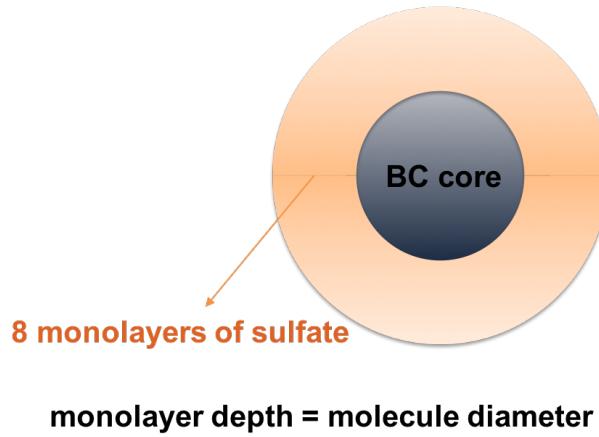


Figure 3.2: Schematic of 8-monolayer of sulfate criterion.

Currently, the condensation of sulfate and ammonia is assumed to be irreversible, whereas the condensation of semi-volatile organics (SOA) can be set to be reversible or irreversible. Sulfate is produced from  $\text{SO}_2$  aqueous oxidation in bulk cloud water by ozone and  $\text{H}_2\text{O}_2$ , based on MOZART treatment (Tie et al. (2001)). An accommodation coefficient (0.65), that is the probability of sticking when the gas molecules encounters the surface of an aerosol particle is used for all the three species (Liu et al. (2012)).

Generally, for each time-step, the model computes the ratio of the condensed soluble species mass in the primary carbon to the mass that is required to age all the particles in that mode based on the monolayer criterion. Then it assumes that the same ratio of the

POM and BC in the primary carbon mode is transferred to the accumulation mode together with the condensed materials. Here we provide a detailed mathematical explanations of this model representation of the condensation scheme.

For each mode  $\textcolor{blue}{k}$ , the rate of gas uptake  $F_{\textcolor{yellow}{n}}$  ( $s^{-1}$ ) is represented as:

$$F_{\textcolor{yellow}{n}} = \int n(\ln D_p) \times Cd(\ln D_p), \quad (3.1)$$

where  $D_p$  is the diameter of the aerosol particles in that mode,  $n(\ln D_p)$  is the lognormal size distribution, and  $C$  is the gas condensation rate (taking sulfuric acid for example):

$$C = 2\pi \times D_p \times V_{\text{diff}} \times F(K_n, A), \quad (3.2)$$

$$V_{\text{diff}} = 0.557 \times 10^4 \times 1.75 \times \frac{T}{P} \quad (3.3)$$

$K_n$  : Knudsen number

$A$  : accommodation coefficient

$F$  : Fuchs-Sutugin correction factor

$V_{\text{diff}}$  : gas diffusivity for  $\text{H}_2\text{SO}_4$

The rate of total gas uptake  $F_{\text{sum}}$  is hence estimated by summing up  $F_n$  over all 4 modes, and the fraction of gas mixing ratio going to each mode  $f_{\textcolor{yellow}{n}}$  is equivalent to the ratio of  $F_{\textcolor{yellow}{n}}$  to  $F_{\text{sum}}$ :

$$F_{\text{sum}} = \sum_{k=1}^4 F_{\textcolor{yellow}{n}}, \quad f_n = \frac{F_{\textcolor{yellow}{n}}}{F_{\text{sum}}}. \quad (3.4)$$

With the above information, the fraction of soluble species condensing on aerosols during  $\Delta t$  is derived as  $(1 - e^{-\Delta t \times F_{\text{sum}}})$ , and the mass mixing ratio of gas uptake that actually going to mode  $\textcolor{yellow}{n}$  during one time-step  $\Delta t$  is:

$$\Delta q_{\textcolor{yellow}{n}} = \textcolor{yellow}{q} \times f_{\textcolor{yellow}{n}} \times (1 - e^{-\Delta t \times F_{\text{sum}}}). \quad (3.5)$$

The model computes the fraction of carbonaceous aerosols being aged  $f_{\text{age}}$  as the ratio of the volume of actually condensed materials  $V_{\text{shell}}$  to the volume of soluble species required to age all the particles in the primary carbon mode  $V_{8-\text{mono}}$ :

$$\begin{aligned} V_{\text{shell}} &= \Delta q_{\text{SO}_4, \text{n}_{\text{pc}}} \times V_{\text{SO}_4} \\ &\quad + \Delta q_{\text{NH}_4, \text{n}_{\text{pc}}} \times V_{\text{NH}_4} \\ &\quad + \Delta q_{\text{SOA}, \text{n}_{\text{pc}}} \times V_{\text{SOA}}, \end{aligned} \quad (3.6)$$

$$V_{8-\text{mono}} = (\pi M_2 / \frac{\pi}{6} M_3) \times d_{8-\text{mono}}, \quad (3.7)$$

$$f_{\text{age}} = \frac{V_{\text{shell}}}{V_{8-\text{mono}}}, \quad (3.8)$$

$V_{\text{SO}_4}$ ,  $V_{\text{NH}_4}$ ,  $V_{\text{SOA}}$ : factors that convert the unit of mixing ratio to  $\text{m}^3/\text{kmol air}$ .

$V_{8-\text{mono}}$ : the volume of sulfate required to age all BC particles.

$V_{\text{core}}$ : the volume of pure BC.

$M_2$ ,  $M_3$ : second and third moment of moment dynamic equation,  $\pi M_2 / \frac{\pi}{6} M_3$  is the ratio of aerosol surface area to volume.

$d_{8-\text{mono}}$ : the thickness of 8 monolayers of sulfate.

The changes of mass in each mode over  $\Delta t$  are then represented as:

$$\Delta q_{n_{\text{pc}}} = \Delta q_{n_{\text{pc}}} - f_{\text{age}} \times q_{n_{\text{pc}}} \quad \text{primary carbon mode}, \quad (3.9)$$

$$\Delta q_{n_{\text{accu}}} = \Delta q_{n_{\text{accu}}} + f_{\text{age}} \times q_{n_{\text{pc}}} \quad \text{accumulation mode}. \quad (3.10)$$

### 3.3 CAM-chem Aging Timescale

In a first-order aging model, the transition of BC mass from fresh to aged mode can be represented by an aging timescale. The model is given by

$$\frac{dM_{\text{fresh}}}{dt} = -\frac{1}{t_{\text{aging}}} M_{\text{fresh}}. \quad (3.11)$$

The mass of freshly emitted BC  $M_{\text{fresh}}$  is the mass of fresh particles. The aging timescales can be represented as the production rate of the inverse of the mass transfer rate and fresh BC mass:

$$t_{\text{aging}} = M_{\text{fresh}} / \left( -\frac{dM_{\text{fresh}}}{dt} \right). \quad (3.12)$$

Any particle transition from the fresh to aged mode during a time step is either by coagulation with other particles or by accumulating a certain amount of condensable materials. So the overall aging timescale  $t_{\text{aging}}$  can be represented as the combination of the aging timescales by condensation  $t_{\text{cond}}$  and by coagulation  $t_{\text{coag}}$ :

$$\frac{1}{t_{\text{aging}}} = \frac{1}{t_{\text{coag}}} + \frac{1}{t_{\text{cond}}}. \quad (3.13)$$

Each of the component aging timescales  $t_{\text{cond}}$  and  $t_{\text{coag}}$  can be derived from the corresponding mass transfer rate by applying Equation 3.12.

In CAM-chem model, BC particles are transferred from the fresh, primary carbon model to the aged, accumulation mode according to mass transfer rates computed from the coagulation and condensation processes respectively. We extracted the mass transfer rates every 6 hours and apply Equation (3.12) and Equation (3.13) to get the corresponding timescales for aging. We would be able to evaluate the contribution of condensation and coagulation to the overall aging by showing their timescales separately. In this study, we evaluated the monthly and annually averaged aging timescales.

### 3.4 PartMC-MOSAIC Aging Timescale

PartMC-MOSAIC (particle resolved model) can track the evolution of individual particles as they evolve through condensation, coagulation, emission and evaporation of SOA (Riemer and West (2013)). It has the advantage in simulating BC aging that no specific criterion assumption is required. Previous study has investigated reduced representation of mixing state for simulating aerosol effects on climate (Fierce et al. (2016)). The mixing timescales that characterized this transformation have been parameterized. In our study, we applied that aging timescale parameterization to our global models outputs to evaluate the accuracy of our global model aging timescales.

Fierce et al. (2016) applied a series of 100 sensitivity scenarios sampled using Latin hypercube sampling (McKay et al. (1979)). Twenty-eight input parameters were varied to represent a range of atmospheric conditions, from highly polluted cases with consequently rapid aging to remote regions with slow aging. Those parameters include environmental variables, aerosol characteristics, aerosol type and gas emissions. In each scenario, they simulate the evolution of carbonaceous particles that is emitted into a population of background aerosol. No new fresh particle will be emitted during this process after the simulation starts, in order to isolate the effects of aging (Fierce et al. (2016)). The normalized error in the number concentration of CCN computed using the particle-resolved composition that track individual particle and the reduced representation of the composition by assuming different mixing states of aerosol population is given by:

$$e_{CCN}(t) = \frac{\int_0^\infty (\tilde{N}_{CCN}(t, s) - N_{CCN}(t, s))ds}{\int_0^\infty \tilde{N}_{CCN,q}(t, s)ds}, \quad (3.14)$$

where  $\tilde{N}_{CCN}$  is the number concentration of CCN using the reduced composition and  $N_{CCN}$  is the CCN using the particle-resolved composition. This error decreases exponentially with time and a regression result has shown a high R-squared value as 93%. The equation can

be expressed as:

$$e_{\text{CCN}}(t) = e_{\text{CCN}}(t_0) \exp\left(-\int_{t_0}^t \frac{1}{\tau_{\text{mix}}(t)} dt\right), \quad (3.15)$$

where  $\tau_{\text{mix}}$  is the overall aging timescale. A particle is either aged by condensation or by coagulation, so the two processes can be treated as separate process and has their own timescale (Fierce et al. (2015)). Accordingly, the overall timescale is found to be a function of the condensation growth rate  $I(t)$  and the particle number concentration  $N(t)$ . So a more specified parameterization of  $\tau_{\text{mix}}$  can be derived by regression of the error on  $I(t)$  and  $N(t)$ :

$$e_{\text{CCN}}(t) = e_{\text{CCN}}(t_0) \exp\left(-k_{\text{cond}} \int_{t_0}^t I(t) dt\right) \exp\left(-k_{\text{coag}} \int_{t_0}^t N(t) dt\right), \quad (3.16)$$

Figure 3.3 is an example of determining  $k_{\text{coag}}$  after setting  $I(t)$  to be 0.

In our study, we computed the number concentration  $N$  as the sum of the particle number concentration of all modes, and the condensation growth rate  $I$  as the volume condensation rate over the total aerosol surface area. As is mentioned in 3.2, the condensation of SOA is reversible, so the total condensation rate can be positive or negative. Since all fresh BC particles are in the primary carbon mode, we computed the condensation growth rate using the volume condensation rate and surface area extracted from the primary carbon mode. The PartMC-MOSAIC parameterized aging timescales can then be represented as:

$$\tau_{\text{overall}} \approx (k_{\text{cond}} I_{\text{cond}} + k_{\text{coag}} N)^{-1}, \quad (3.17)$$

Fierce et al. (2016) has shown the estimated e-folding time  $\tau_{\text{overall}}$  computed from the approximate range of condensation growth rate and number concentration for specific locations (Figure 3.4). This range can also be used as a reference for global models.

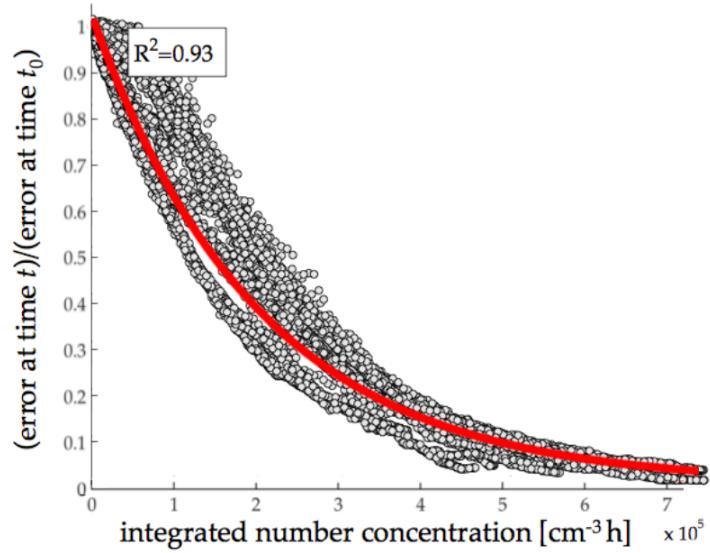


Figure 3.3: Raw data that is used to construct the regression function (grey dots) and the resulting regression function (black line) for simulations including only coagulation, without condensation (Fierce et al. (2016)).

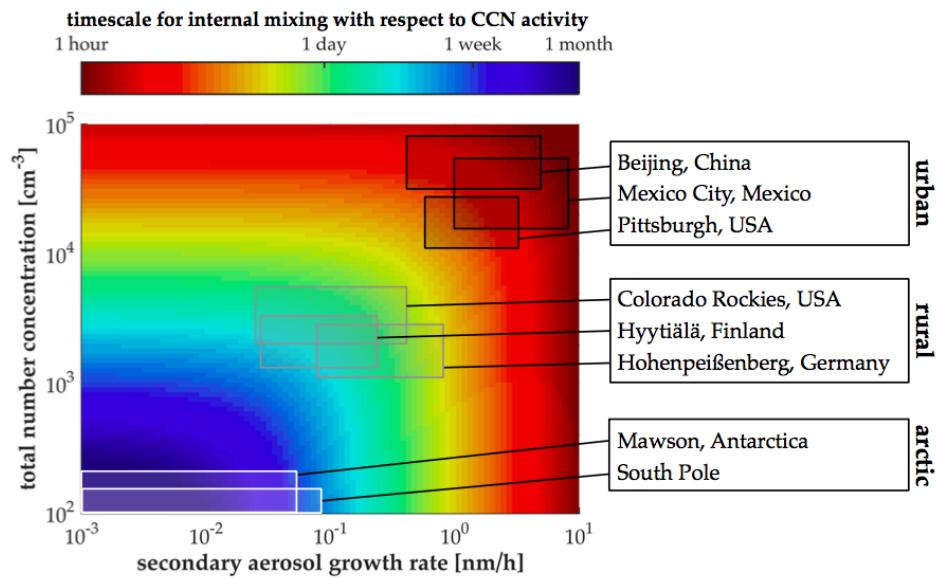


Figure 3.4: Estimated e-folding time for the error in CCN activity from the internal mixture approximation. (Fierce et al. (2016)).

# Chapter 4

## Results

### 4.1 BC Emissions

We use the Intergovernmental Panel on Climate Change (IPCC) Fifth Assessment report (AR5) gridded POM and BC emissions for the period 1850-2010 in decadal increments (Lamarque et al. (2010)). Sources include anthropogenic emissions originating from domestic, energy, industry, transportation, waste treatment and ship activity sectors, and natural emissions (elevated) from forest fire and grass fire. The distribution of anthropogenic BC emissions (Figure 4.2) shows ocean going ship tracks in the Arctic region, and the relatively high emission fluxes in Europe might also be a regional contribution to the BC burden in the Arctic. BC emission fluxes are maximum in industrial regions such as Southeast Asia and biomass burning regions such as Southern Africa and Amazon rainforest. A strong seasonal variation in natural BC emissions can be observed (Figure 4.1), where the fluxes in September are more than 1000 times higher than the fluxes in March for regions like Amazon rainforest and South Africa, and 10 times higher for the red-colored regions (see left-bottom panel in Figure 4.1) throughout Russia and Europe.

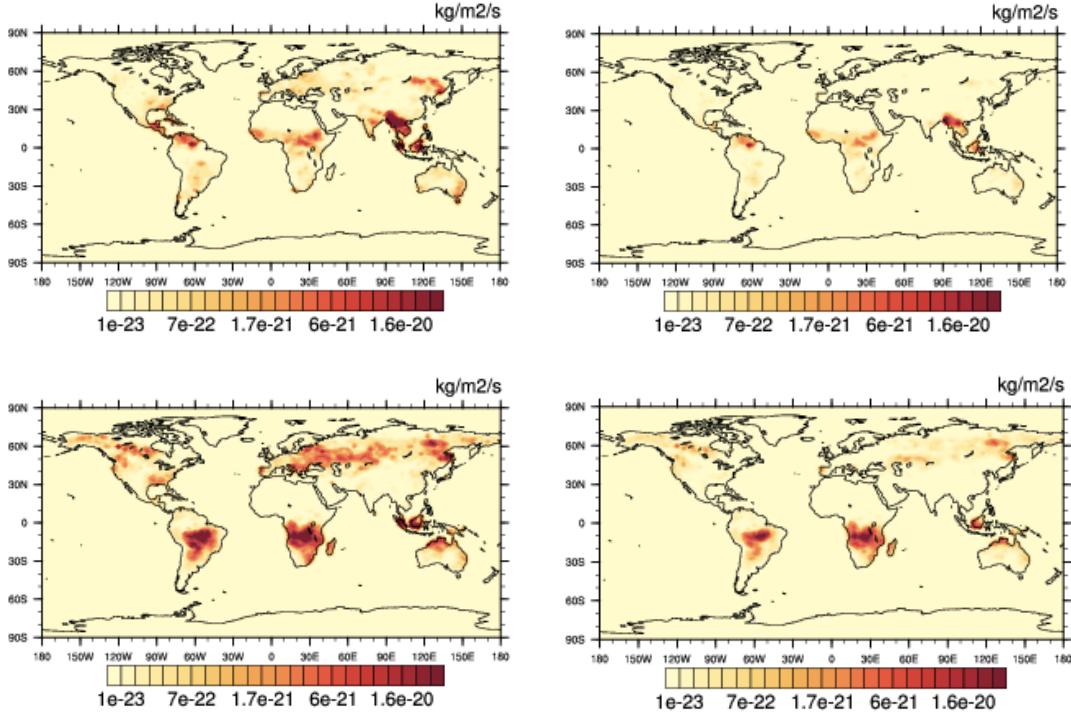


Figure 4.1: Monthly mean forest fire and grass fire fluxes of BC for March (top panels) and September (bottom panels), at heights of 0–126 m (left panels) and 278–454 m (right panels).

## 4.2 Sensitivity of BC Burden and ~~Climate Effect to the Number of Monolayers~~

To explore the model sensitivity to the ~~number of monolayer of sulfate condensation criterion~~, we conducted four experiments at  $1.9^\circ \times 2.5^\circ$  from January 1 2010 to December 31 2010 with offline meteorology, and set the number of mono-layers to 1, 2, 4 and 8 (default), respectively, represented as L1, L2, L4 and L8. For a larger number of monolayers criterion, more coating material is required to transfer the carbonaceous aerosols from the primary carbon mode to the accumulation mode, hence BC will stay longer in the primary carbon mode and the aging rates will be slower. Consequently, the BC concentration will be higher because less BC is transferred to the hydrophilic, accumulation mode that is subject to wet deposition.

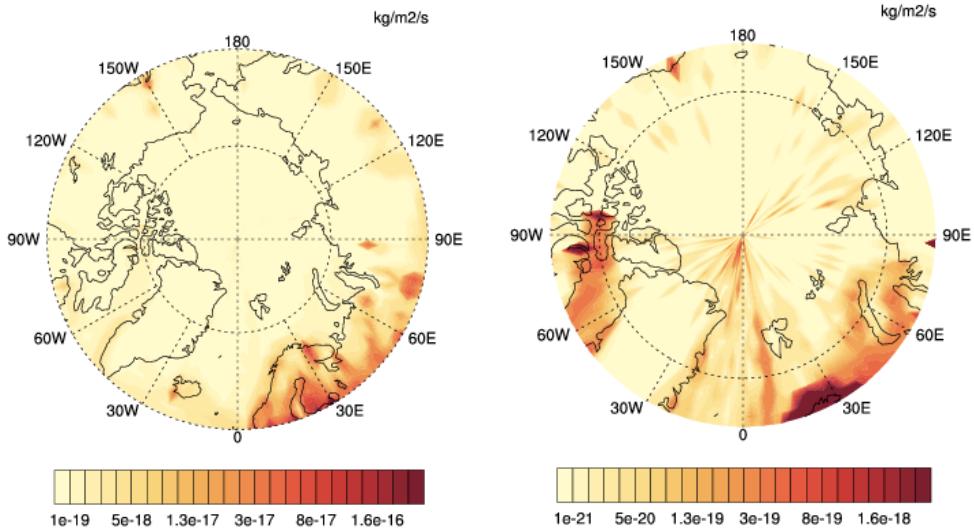


Figure 4.2: Annual mean anthropogenic emissions of BC emitted at the surface for the year 2010 for 60–90 °N (left) and 70–90 °N (right).

#### 4.2.1 Horizontal Profiles of BC burdens

Figure 4.3 shows the horizontal distributions of BC mixing ratio near the surface in the L8 case (top panel), and the relative differences between L1 and L8 cases (bottom panel), denoted by  $(L1 - L8)/L8$ . Generally, BC concentrations in the L1 case are lower than L8 case throughout the globe with maximum differences in the annually averaged BC mixing ratio of 16% near the surface. This is because the aging is faster with a smaller threshold of the number of monolayers, and hence more fresh BC will be transferred to accumulation mode. The accumulation mode aerosols are then subject to wet removal, which will prevent them from traveling to remote regions.

We observed that the relative differences increase along the pathway of BC as it is transported away from the source regions, such as the South Atlantic Ocean downwind of South Africa or the North Pacific near Southeast Asia, both show higher differences than their surrounding regions and can be matched to obvious BC burden gradient along BC pathways over the ocean. Simulated BC burden is most sensitive to the choices of aging criterion in the high-latitude regions where the background concentrations are low, with

maximum differences in the annually averaged BC mixing ratio of 16% near the surface.

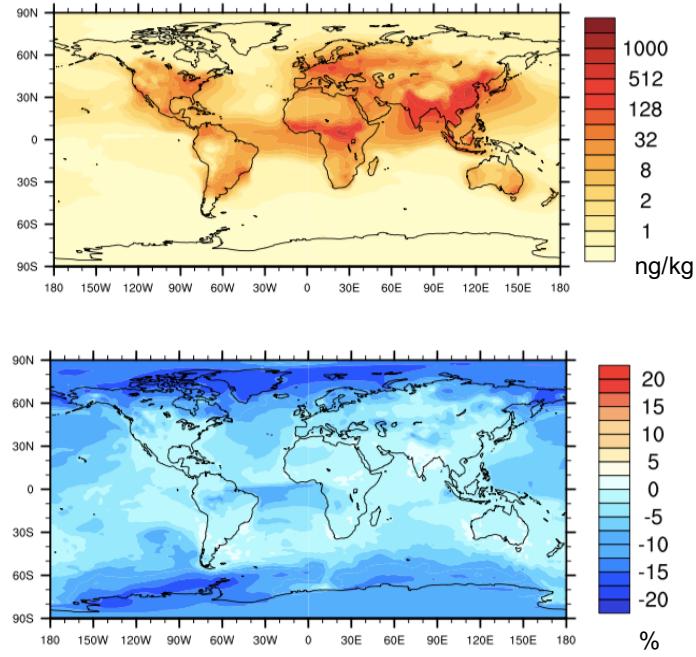


Figure 4.3: Annual BC mass mixing ratio (top) and relative differences denoted by  $(L_1 - L_8)/L_8$  (bottom).

#### 4.2.2 Vertical Profiles of BC burdens

Figure 4.6 and 4.7 show the sensitivity of BC vertical distributions to the number of monolayers in six regions (Figure 4.5). Region 1 and 2 are distant from emission sources while the other four regions are dominated either by biomass burning activities (region 3 and 4) or by emissions from both anthropogenic and biomass burning activities (region 5 and 6). It is obvious that BC distributions in distant regions are more sensitive to the number of monolayers (region 1 and 2 compared to 3–6). The maximum relative differences among the six regions are found at 400 hPa in region 2 during March, indicating that the highest model sensitivity can appear at high altitudes near the Arctic. Throughout the source regions, a decreasing trend of BC burdens with height is shown whereas an increasing trend is observed instead for distant regions, probably due to the wet removal process that prevents

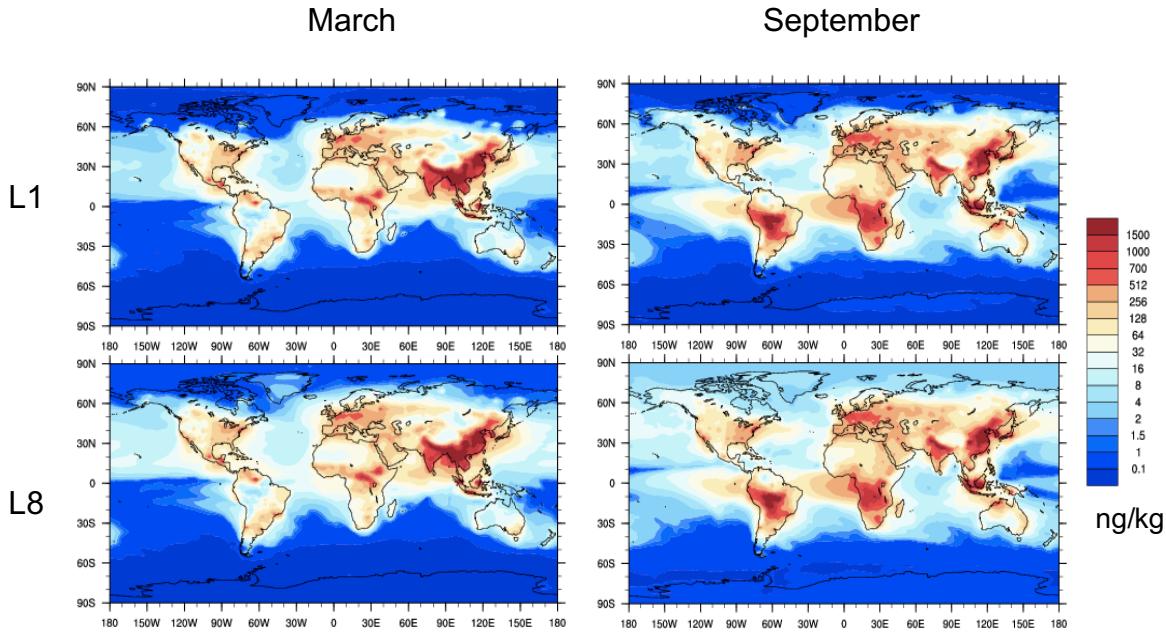


Figure 4.4: Horizontal plots of BC concentrations near the surface for different seasons (March and September).

lower level BC from being transported far away from its sources. These results are still to be further evaluated in the future, considering that the model simulations tend to overestimate BC burdens in the upper troposphere and underestimate BC burdens at altitudes below 400 hPa (Liu et al. (2016)).

Figure 4.6 and 4.7 also compare the model simulated BC vertical profiles in March and September in order to explore their seasonal variability. Regions dominated by biomass burning have significant increases of BC burdens in September by **10 order of magnitude** (region 3 and 4). BC **concentrations** tend to be maximum in September because of South American and Africa biomass burning activities in the dry season (Liu et al. (2016)). The poleward transport can also be seen from the vertical profile of region 1 at high altitudes above 400 hPa, especially for September. The differences among the four MAM4 experiments are smaller in September especially in the Arctic and oceanic regions. In addition, more BC are concentrated in the upper troposphere in September, probably because of the biomass

burning activities produces BC at higher altitudes compared to other sources.

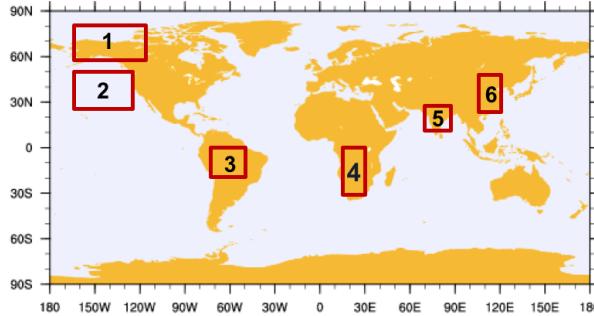


Figure 4.5: Horizontal map of six regions: Region 1 ( $162^{\circ}\text{W}$ – $112^{\circ}\text{W}$ ,  $60^{\circ}\text{N}$ – $80^{\circ}\text{N}$ ), Region 2 ( $160^{\circ}\text{W}$ – $130^{\circ}\text{W}$ ,  $30^{\circ}\text{N}$ – $50^{\circ}\text{N}$ ), Region 3 ( $75^{\circ}\text{W}$ – $55^{\circ}\text{W}$ ,  $10^{\circ}\text{S}$ – $1^{\circ}\text{N}$ ), Region 4 ( $15^{\circ}\text{E}$ – $30^{\circ}\text{E}$ ,  $30^{\circ}\text{S}$ – $0^{\circ}\text{N}$ ), Region 5 ( $73^{\circ}\text{E}$ – $83^{\circ}\text{E}$ ,  $15^{\circ}\text{N}$ – $27^{\circ}\text{N}$ ), Region 6 ( $108^{\circ}\text{E}$ – $120^{\circ}\text{E}$ ,  $23^{\circ}\text{N}$ – $43^{\circ}\text{N}$ ).

Figure 4.8 shows the horizontal plots of regional BC concentrations at 859 hPa for the arctic region. The patterns of BC distributions in the two MAM4 experiments are quite different, in addition to their regional mean differences among the cases that has been explained in the previous paragraphs. This finding indicates that the sensitivity of not only the magnitude of BC burdens, but also their latitudinal and longitudinal patterns are both very sensitive to the aging criterion, especially in the middle and upper troposphere. So understanding the BC aging process so as to improve the reliability of the number-of-monolayer criterion is quite important for the accuracy of CAM-chem MAM4 BC simulations.

### 4.2.3 BC Deposition Flux

We also explored the sensitivity of BC deposition flux through both wet and dry processes to the aging criterions, and compared their differences between two cases (L1 v.s. L8). Figure 4.9 and 4.10 shows the relative differences of the monthly averaged wet deposition rates and dry deposition rates respectively, and compares the seasonal differences.

As is expected, the wet deposition rates with a smaller threshold of coating thickness giving to faster aging rates are higher in the main source regions (South East Asia, Europe,

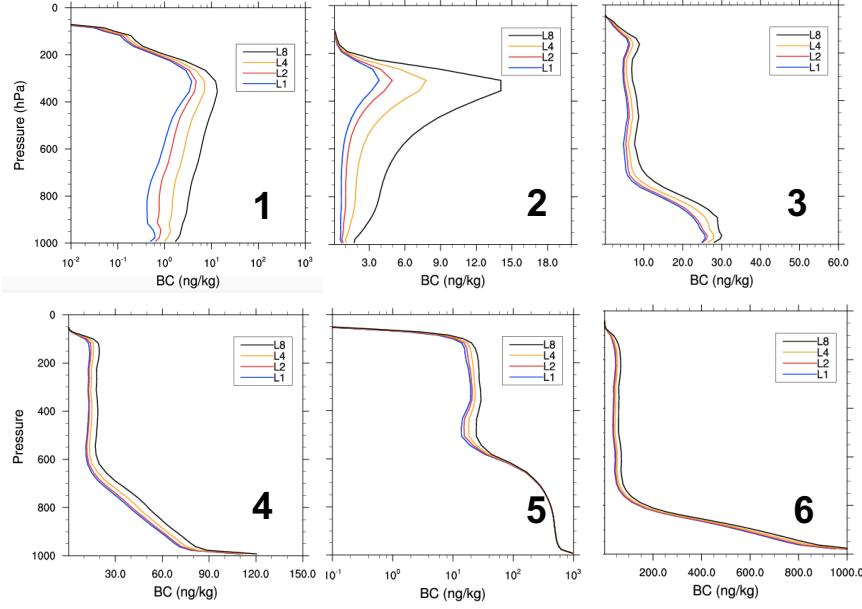


Figure 4.6: Vertical profiles of mean burdens for six regions corresponding to Figure 4.5 in March, 2010.

South America, Africa, etc.) by a factor around 0.4. The wet deposition rates in case L1 become smaller than case L8 in far regions such as the distant oceans and polar regions, with the highest relative differences attaining 90 % in the Arctic. This is because BC stays longer in the primary carbon mode with a larger number of monolayer criterion, and hence can be transported to distant regions. We also noticed the high values at the north edge of mainland Russia or at the south edge of South America continent, primarily because of the small values of the background concentrations (the denominator of the relative difference equation) ~~and thus are not much concerned in our study~~.

The dry deposition rates are positively related to the overall number concentration of BC aerosols (Figure 4.4), where higher concentrations lead to larger dry deposition fluxes. So similarly to our analysis in section 4.2, BC concentrations in L1 case are lower than L8 case throughout the globe, and the relative differences increase along the pathway of BC as it is transported away from the source regions. The dry scavenging rates ~~is~~ most sensitive to the choices of aging criterion in the high-latitude regions, with maximum differences of

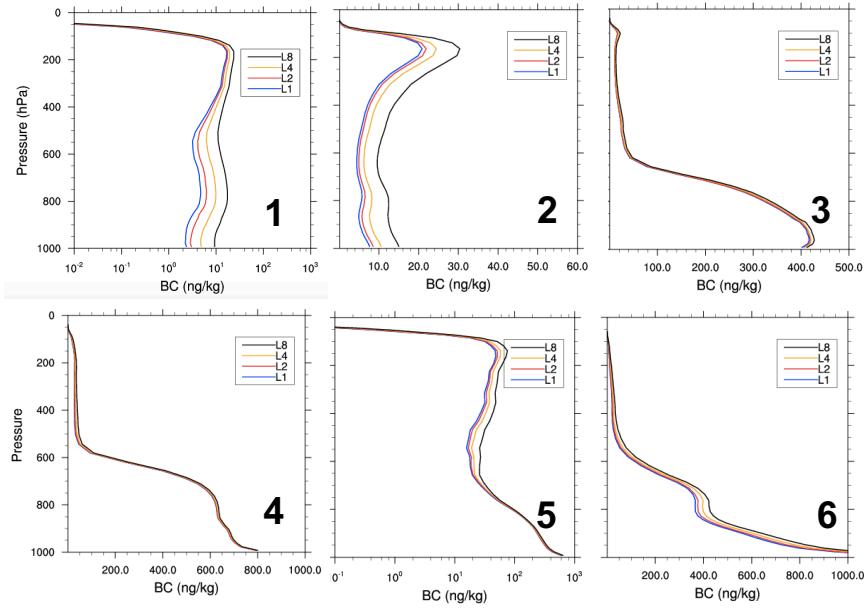


Figure 4.7: Vertical profiles of mean BC burdens for six regions corresponding to Figure 4.5 in September, 2010.

99% in the Arctic for September.

#### 4.2.4 ~~Monolayer Effect on Radiative Forcing~~

We analyzed the sensitivity of BC radiative forcing to the aging criterions, and compared their differences between L1 and L8. Figure 4.12 and 4.11 shows the horizontal plots and polar plots of the annually averaged direct radiative forcing of black carbon aerosols respectively, and compares the differences between 1 monolayer and 8 monolayer as its condensation criterion.

A smaller threshold of coating thickness gives rise to faster aging rates and a larger portion BC in the accumulation mode (internally mixed). The accumulation mode aerosols are subject to wet scavenging and removed as they are transported to distant regions. We have shown in the previous sections that generally BC mixing ratios in the L1 case are lower than L8 case throughout the globe, and the simulated BC burden is most sensitive to the choices of its aging criterion at high latitudes where the background concentrations are low.

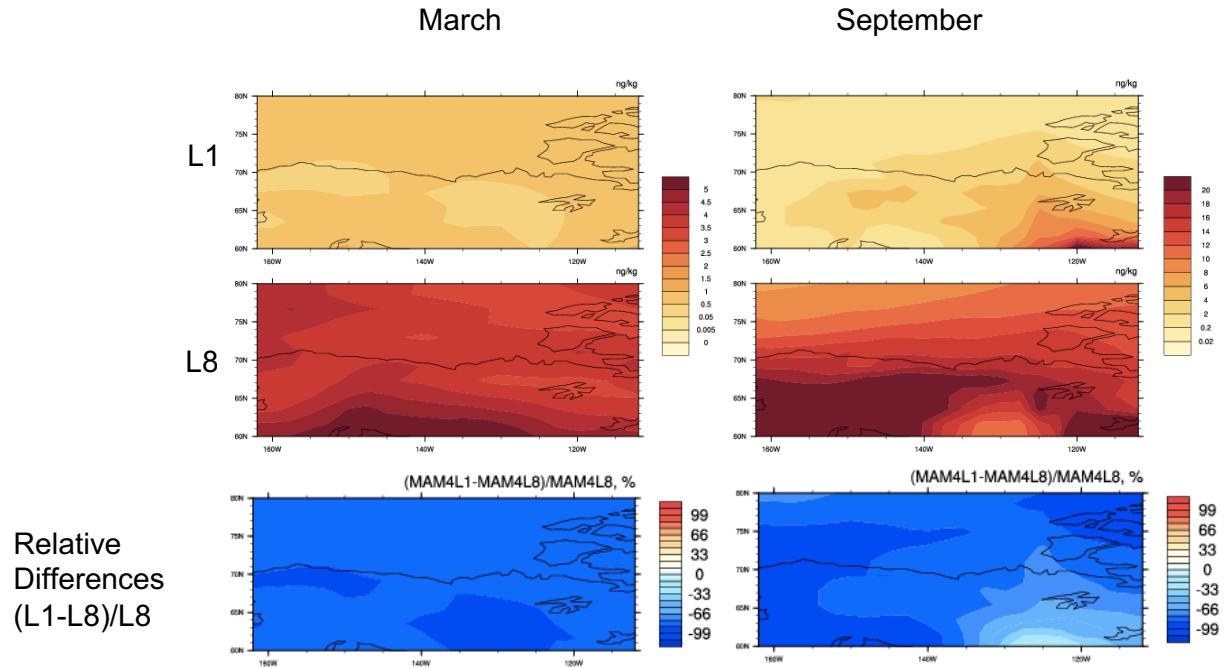


Figure 4.8: Horizontal plots of regional BC concentrations at 859 hPa covering region 1 in case L1 (top) and L8 (middle), and the relative differences (bottom) denoted by  $(L1 - L8)/L8$ .

Consequently, the magnitude of the spatially averaged radiative forcing is  $0.18 \text{ W}\cdot\text{m}^{-2}$  in the L1 case, lower than the radiative forcing of  $0.48 \text{ W}\cdot\text{m}^{-2}$  in the L8 case because less BC are there with larger number of monolayers, and consequently faster aging and wet deposition. The magnitudes are higher in the main source regions (South East Asia, Europe, South America, Africa, etc.) with a maximum of around  $15 \text{ W}\cdot\text{m}^{-2}$  in both cases.

## 4.3 Mixing State of Species in MAM4

### 4.3.1 Volume Fraction of Species in the Primary Carbon Mode and Accumulation Mode

Figure 3.1 shows a schematic of the species component in MAM4. In this section, we will discuss the volume fraction of each species in the accumulation mode and primary carbon

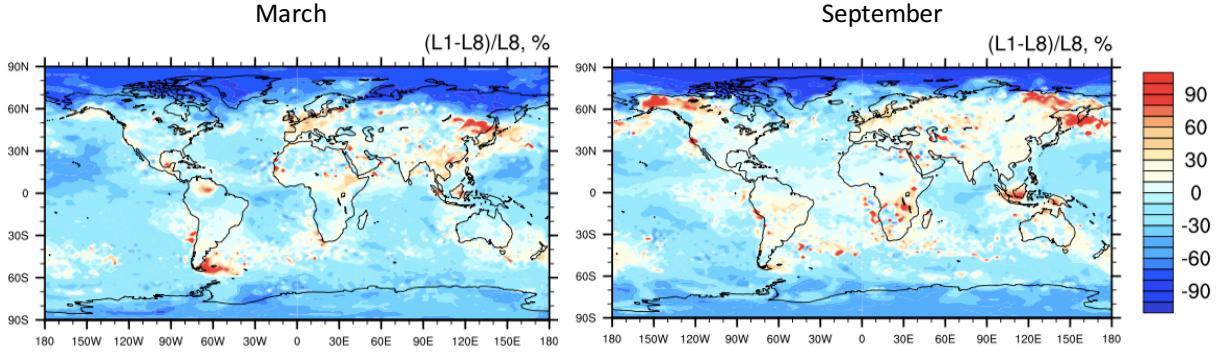


Figure 4.9: Horizontal plots of the relative differences in BC wet deposition flux with different aging criterion (L1 v.s. L8) for March (left) and September (right).

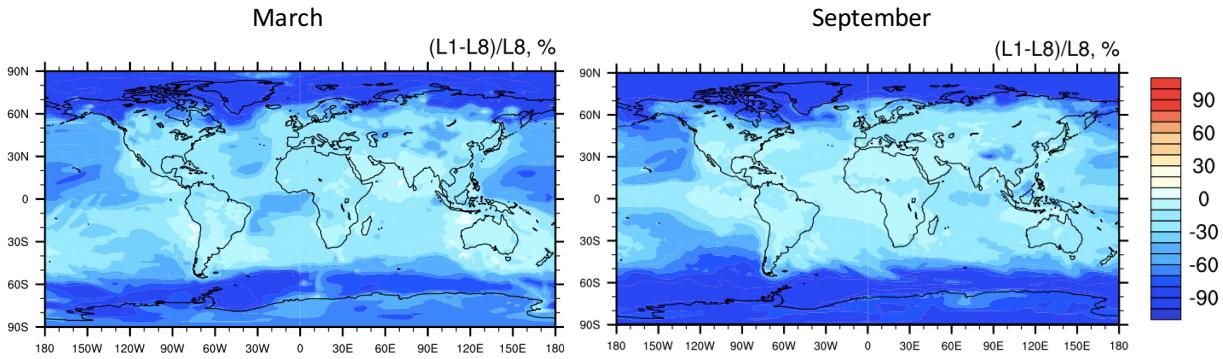


Figure 4.10: Horizontal plots of the relative differences in BC dry deposition flux with different aging criterion (L1 v.s. L8) for March (left) and September (right).

mode respectively, as an indication to the species composition abundance and mixing states in both modes. We focus on the two modes because they both contain carbonaceous aerosols. More specifically, BC-containing aerosols tend to be internally mixed with Sulfate and SOA in the accumulation mode and ~~be~~ externally mixed in the primary carbon mode.

Figure 4.13 and 4.14 illustrate the volume fraction of the six species in the accumulation mode. SOA takes up more than 70% of the total volume over the tropical and subtropical continents including South America, South Africa, South Asia and Australia. This ~~may~~ indicates that SOA plays a dominating role in BC aging through condensation over those regions (we will explain ~~it~~ in the last section). Sulfate has high volume fractions (> 60%)

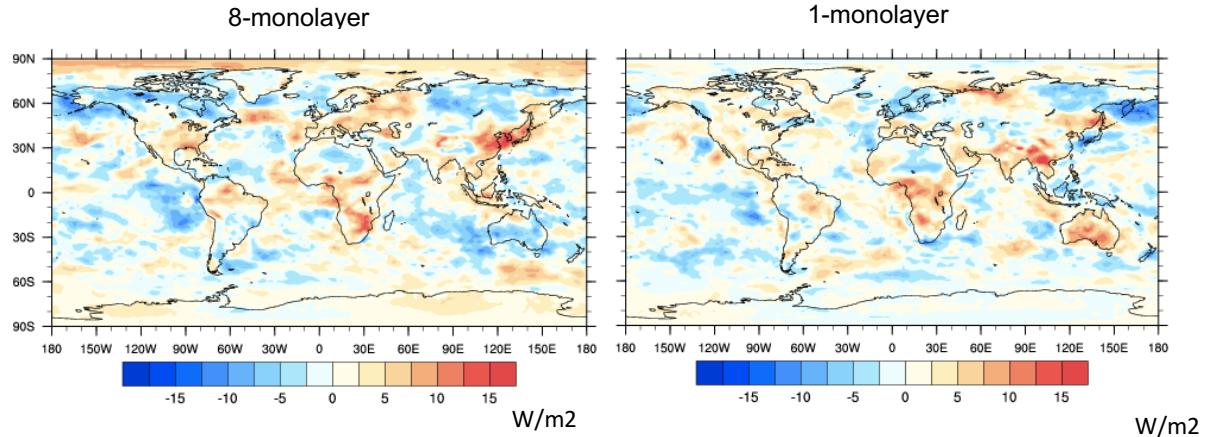


Figure 4.11: Horizontal plots of the direct radiative forcing of BC aerosols with different aging criterion (L1 v.s. L8) near the surface (992hPa).

at polar regions, primarily because its formation by homogeneous nucleation in a system of binary  $\text{H}_2\text{SO}_4 - \text{H}_2\text{O}$  mixture is more likely at a lower ambient temperatures (e.g, Hamill et al. (1977); Yue and Deepak (1982)), and the bulk volume of all species at high latitudes tend to be low. In addition, sulfur species that ultimately forms sulfate can be ejected directly into the stratosphere through volcanic eruptions or the diffusion of sulfide from the troposphere (e.g., Castleman Jr et al. (1974)). So that stratospheric portion can be protected from the tropospheric wet deposition and will be able to travel to the distant polar regions. In a later section, we will show that the condensation of SOA and sulfate plays a dominating role in BC aging. We also noticed that BC contribute a small volume fraction (< 5%) in the accumulation mode. Sea salt volume fractions are highest over the ocean (> 90%) whose values decrease with height. The volume fraction of BC and OC are roughly proportional to each other, indicating their co-emission and internally mixing with each other. For the primary carbon mode (Figure 4.15), OC takes up the major portion.

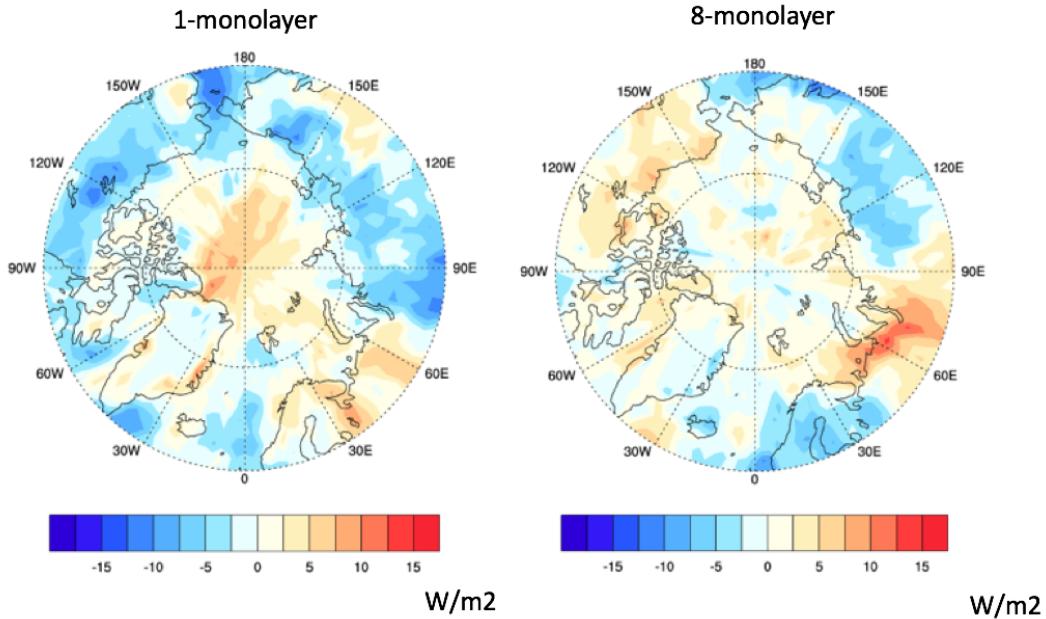


Figure 4.12: Polar plots of the direct radiative forcing of BC aerosols with different aging criterion (L1 v.s. L8) near the surface (992hPa).

### 4.3.2 BC Mixing States Sensitivity to the Aging Criterion

The mixing state of atmospheric BC particles with those hydrophilic aerosol components can affect both their CCN/IN activities and radiative properties. Figure 4.16 shows the comparison of BC mixing states between two MAM4 experiments, represented by the fraction of BC mixing ratio in the accumulation mode. We noticed that the model in the L1 experiment has more than 90 % of BC mass in the accumulation mode almost everywhere throughout the globe. The fraction is even less than 10 % for some regions in the Arctic (Greenland, Barents sea, etc.). Most of BC is in primary carbon mode are externally mixed in the Arctic that should be of our interest, considering the fact that BC has a strong climate effect in the Arctic, and its mixing states will affect its climate forcing properties such as CCN properties and optical depth properties.

We have found that the BC mixing ratios in the L1 case are lower than the L8 case throughout the globe, and the simulated BC burden is most sensitive to the choices of its

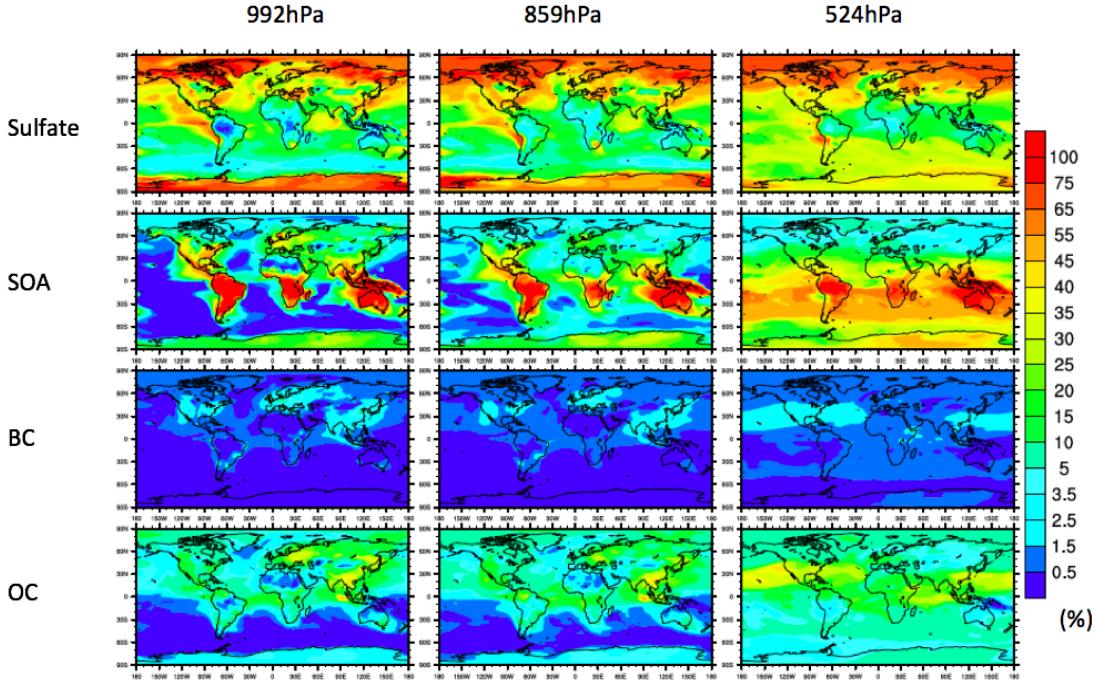


Figure 4.13: Annual volume fraction of sulfate, secondary organic aerosol (SOA), black carbon (BC) and organic carbon (OC) near the surface (992hPa) in the accumulation mode.

aging criterion at high latitudes. This is consistent with our results that the mixing level is more sensitive to the aging criterion at high latitudes. Comparing L8 to the L1 case, the high fraction of BC in the primary carbon mode at high latitudes seems a bit confusing because we expect BC to be more internally mixed as it is transported to distant regions. It can be understood that less BC is aged in L8 case, so relatively a larger portion of BC in the L8 accumulation mode is deposited during its transport and little is left in the accumulation mode when the population arrives the Arctic. The spatial sensitivity of BC mixing level to latitudes is more intense in September, considering there is stronger seasonal precipitation in NH Summer (e.g., monsoon season).

In the next section, we will also discuss whether the SP2 observed BC can be representative of the total BC especially in the Arctic region. We will show that most of BC detected by SP2 measurements is also externally mixed in the Arctic, and SP2 can capture most of BC in accumulation mode (internally mixed) when it is close to the source regions.

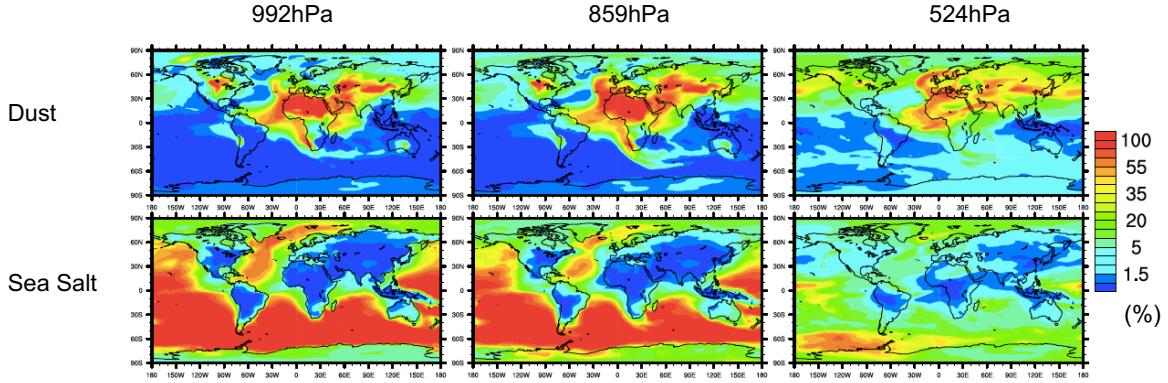


Figure 4.14: Annual volume fraction of dust and sea salt (SST) near the surface (992hPa) in the accumulation mode.

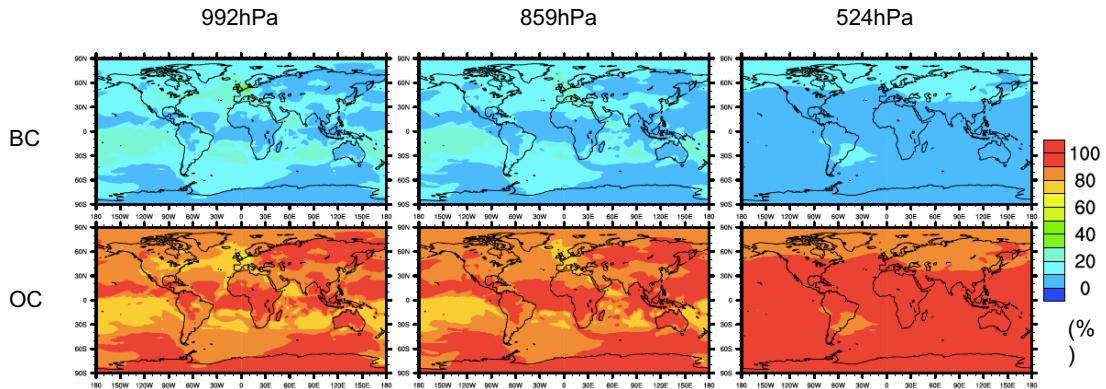


Figure 4.15: Annual volume fraction of black carbon (BC) and organic carbon (OC) near the surface (992hPa) in the primary carbon mode.

## 4.4 Mass Fraction and Mixing States Observed by SP2 measurement

The SP2 instrument measures the BC particle cores over a calibrated volume equivalent diameter (VED) range of 55–400 nm, which is unlikely to represent the total ambient number and mass concentrations of BC particles (Reddington et al., 2013). In order to compare CAMChem model simulated BC with observations, we estimated the mass fraction of modeled BC in the size range corresponding to SP2 measurement. SP2 number-detection

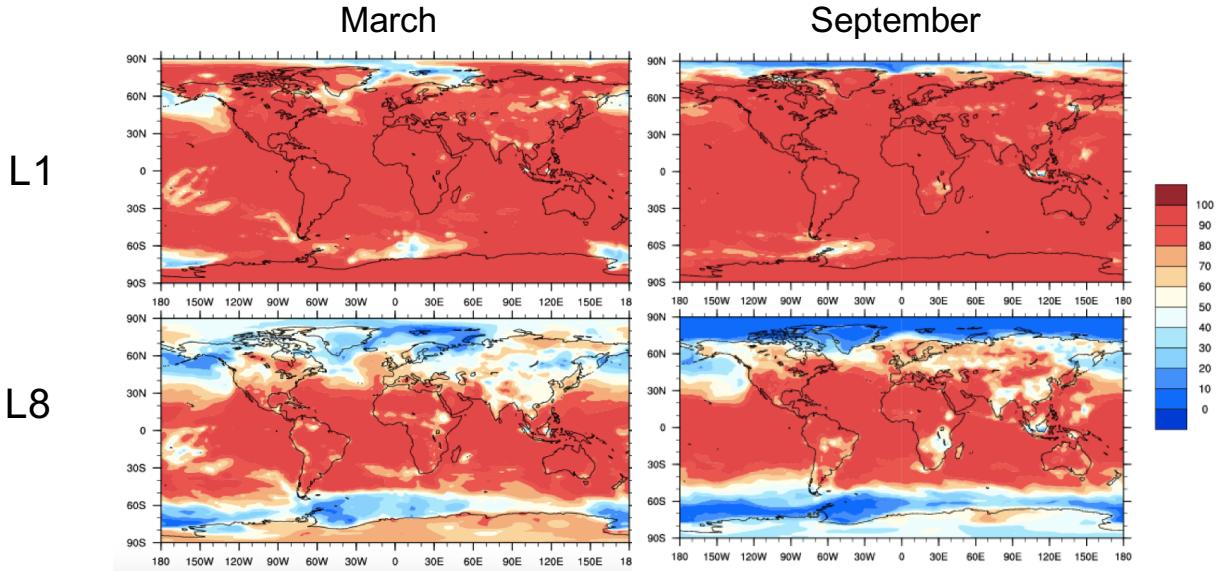


Figure 4.16: Fraction of BC mixing ratio in the accumulation mode to total BC mixing ratio (in %) for L1 case (top panels) and L8 case (bottom panels) in March (left panels) and September (right panels).

efficiency at sea level pressure is reported to be 100% for BC above 90 nm VED (Schwarz et al., 2010), so following Reddington et al. (2013), we use 90 nm–400 nm as the efficient diameter range of SP2 measurement in this study.

#### 4.4.1 BC Core Diameter

MAM4 assumes lognormal distribution for each mode. Before we can estimate the volume fraction of BC within the SP2 measurement size range, we first derive the geometric mean diameter of BC core.

The mean diameter of BC core can be calculated as:

$$d_g = (d_{\text{mixed}}^3 \times f_{\text{BC}})^{\frac{1}{3}},$$

where  $d$  is the geometric mean diameter of BC core,  $d_{\text{mixed}}$  is the geometric mean diameter of internally mixed particles (**extracted from model**), and  $f_{\text{BC}}$  is the volume fraction

of BC in accumulation mode. For the following estimation of volume fraction within SP2 measurement size range, we refer to the geometric mean diameter of BC particles as its core diameter.

#### 4.4.2 Volume Fraction Within the Size Range of SP2

##### Measurement

The CDF of lognormal number distribution of the BC cores in the diameter range between  $d_1$  and  $d_2$  is:

$$N(d_1, d_2) = \frac{1}{\ln\sigma_g\sqrt{2\pi}} \int_{d_1}^{d_2} e^{-\frac{(\ln d - \ln d_g)^2}{2\ln^2\sigma_g}} d(\ln d),$$

where  $d_g$  is the geometric mean diameter of the BC core distribution (extracted from the model, varying temporally and spatially).

The third moment of the lognormal distribution of BC core between 90 and 400 nm  $M_3(d_1, d_2)$  (proportional to the volume of BC cores whose diameter is within that size range) is:

$$\begin{aligned} M_3(d_1, d_2) &= \frac{1}{\ln\sigma_g\sqrt{2\pi}} \int_{d_1}^{d_2} d^3 e^{-\frac{(\ln d - \ln d_g)^2}{2\ln^2\sigma_g}} d(\ln d) \\ &= \frac{e^{\frac{k^2}{2}\ln^2\sigma_g + k\ln d_g}}{\ln\sigma_g\sqrt{2\pi}} \int_{d_1}^{d_2} d^3 e^{-\frac{(\ln d - \ln d_{gv})^2}{2\ln^2\sigma_g}} d(\ln d), \end{aligned}$$

where the geometric mean diameter of BC volume is represented as  $\ln d_{gv} = \ln d_g + 3\ln\sigma_g$

So the mass fraction of the BC cores in the size range between  $d_1$  and  $d_2$  (in each mode) is derived as:

$$\begin{aligned}
F(d_1, d_2) &= \frac{\frac{1}{\ln\sigma_g\sqrt{2\pi}} \int_{d_1}^{d_2} d^3 e^{-\frac{(\ln d - \ln d_g)^2}{2\ln^2\sigma_g}} d(\ln d)}{\frac{1}{\ln\sigma_g\sqrt{2\pi}} \int_{-\infty}^{+\infty} d^3 e^{-\frac{(\ln d - \ln d_g)^2}{2\ln^2\sigma_g}} d(\ln d)} \\
&= \frac{\frac{e^{\frac{k^2}{2}\ln^2\sigma_g + k\ln d_g}}{\ln\sigma_g\sqrt{2\pi}} \int_{d_1}^{d_2} e^{-\frac{(\ln d - \ln d_{gv})^2}{2\ln^2\sigma_g}} d(\ln d)}{\frac{e^{\frac{k^2}{2}\ln^2\sigma_g + k\ln d_g}}{\ln\sigma_g\sqrt{2\pi}} \int_{-\infty}^{+\infty} e^{-\frac{(\ln d - \ln d_{gv})^2}{2\ln^2\sigma_g}} d(\ln d)} \\
&= \frac{1}{\ln\sigma_g\sqrt{2\pi}} \int_{d_1}^{d_2} e^{-\frac{(\ln d - \ln d_{gv})^2}{2\ln^2\sigma_g}} d(\ln d) \\
&= \frac{1}{2} [\operatorname{erf}(\frac{\ln d_2 - \ln d_{gv}}{\sqrt{2}\ln\sigma_g}) - \operatorname{erf}(\frac{\ln d_1 - \ln d_{gv}}{\sqrt{2}\ln\sigma_g})],
\end{aligned}$$

The above mass fraction for primary carbon mode ( $F_{pc}(d_1, d_2)$ ) and for accumulation mode ( $F_{accu}(d_1, d_2)$ ) do not add up to 1:

$$F_{accu}(d_1, d_2) + F_{pc}(d_1, d_2) \neq 1.$$

**Within the size range (90–400 nm),** the ratios of the mass of BC cores in each mode to the total mass of BC cores are computed as:

$$\begin{aligned}
f_{accu} &= \frac{F_{accu}(d_1, d_2)M_{accu}}{F_{accu}(d_1, d_2)M_{accu} + F_{pc}(d_1, d_2)M_{pc}}, \\
f_{pc} &= \frac{F_{pc}(d_1, d_2)M_{pc}}{F_{accu}(d_1, d_2)M_{accu} + F_{pc}(d_1, d_2)M_{pc}},
\end{aligned}$$

$$f_{accu} + f_{pc} = 1,$$

where  $f_{accu}$  is the mass fraction of BC cores in accumulation mode,  $f_{pc}$  is the mass fraction of BC cores in primary carbon mode,  $M_{accu}$  and  $M_{pc}$  are the mass mixing ratio of BC cores in accumulation mode and primary carbon mode, respectively.

Sketch of F and f are shown in Figure 4.17, Figure 4.18 and Figure 4.19.

Generally, the mixing ratio of BC particles in the accumulation mode ( $M_{accu, BC}$ ) is higher than that in the primary carbon mode ( $M_{pc, BC}$ ) when it is distant from the source regions

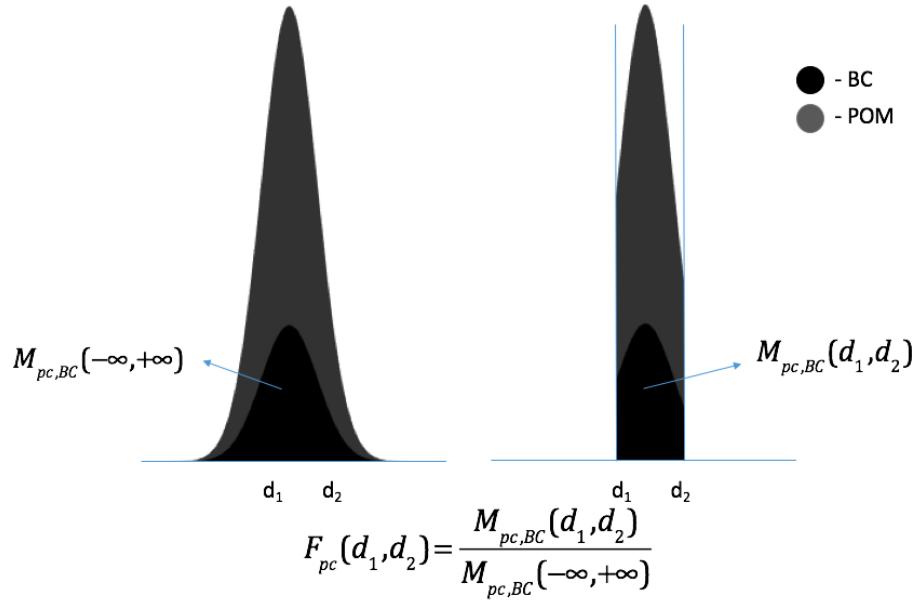


Figure 4.17: BC mass fraction within the SP2 size range for primary carbon mode  $F_{pc}(d_1, d_2)$ .

(e.g., Indian Ocean, Atlantic Ocean) (Figure 4.20), primarily because black carbon will be transferred from the fresh, hydrophobic mode (primary carbon) into the aged, hydrophilic mode (accumulation) during transportation. We also observe that for some Arctic regions (e.g., Greenland), however, BC mixing ratio can be higher in the primary carbon mode than in the accumulation mode.

BC mass fractions within the SP2 size range ( $F_{pc}(90 \text{ nm}, 400 \text{ nm})$  and  $F_{accu}(90 \text{ nm}, 400 \text{ nm})$ ) are shown in Figure 4.21. For March (upper left panel), SP2 would be able to detect most of BC (>60%) over the ocean and throughout the North Pole in primary carbon mode, and less of BC (<40%) in regions like central Africa, Northeast Asia and Australia. This portion is higher (lower left panel) when it is close to the sources (e.g., South East Asia, Europe), and lower at high latitudes in accumulation mode. For September (upper right panel), however, SP2 can only detect less than >20% of BC in the Arctic in both modes. We also observe that the distributions of the above mass fractions (Figure 4.21) have similar pattern with the size distributions of BC cores (Figure 4.22), which indicates that the low values of the

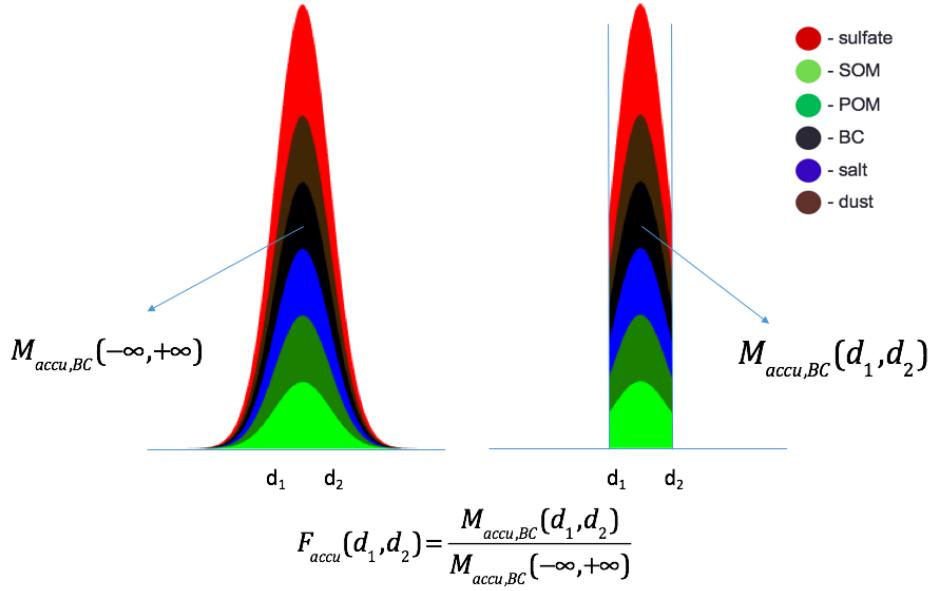


Figure 4.18: BC mass fraction within the SP2 size range for accumulation mode  $F_{accu}(d_1, d_2)$ .

mass fraction ( $F_{pc}(90 \text{ nm}, 400 \text{ nm})$  and  $F_{accu}(90 \text{ nm}, 400 \text{ nm})$ ) are highly due to the small geometric mean diameter of BC cores ( $<90 \text{ nm}$ ) in that region.

The ratios of BC mixing ratio in each mode within the SP2 size range to the total BC mixing ratio within the SP2 size range ( $f_{pc}$  and  $f_{accu}$ ) are shown in Figure 4.23, which represents the mixing states of BC particles that are captured by SP2 measurements. In March, most of BC aerosols are internally mixed at low and middle latitudes ( $f_{accu}>60\%$ ), whereas more of them are externally mixed at high latitudes ( $f_{pc}>60\%$ ). In September, the mixing state can vary spatially in the Arctic region. The above results should be taken into consideration when comparing modeled BC with SP2 measurements. Fractions in top and bottom panels should add up to 1.

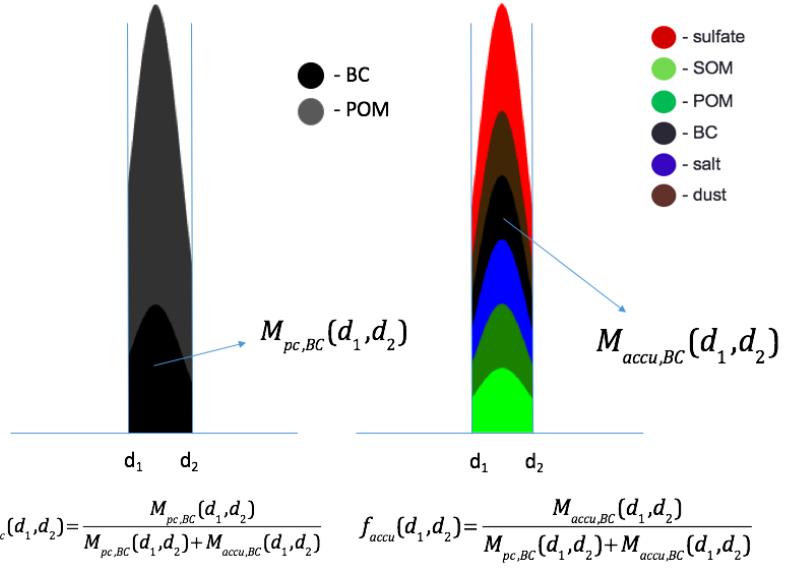


Figure 4.19: The ratios of BC mass in the SP2 size range in each mode to the total BC mass in the SP2 size range, for primary carbon mode  $f_{pc}(d_1, d_2)$  and for accumulation mode  $f_{accu}(d_1, d_2)$ .

## 4.5 BC Mass Fraction Activated by Coating Material

CAM-chem MAM4 assumes that the hydrophobic BC particles in primary carbon mode will be activated and transferred to the accumulation mode after condensing a certain number of sulfate molecular layers, and it uses 8-monolayer as default. The size and hygroscopicity of BC aerosols will be increased as more condensable material is coated onto their surface, and this will lead to a decreasing supersaturation ratio to activate the particles. In MAM4, more coating material is required to age BC for larger number of monolayers, so the critical supersaturation ratio will be decreases for a larger number of monolayers. Liu et al. (2012) has estimated the critical supersaturation for 3 and 8 monolayers of sulfate to produce CCN from BC aerosols, and the values are 0.49% and 0.32% respectively.

Following Liu at al. (2012), we derived similar results and the values are 0.43% and 0.28% for 3 and 8 monolayers respectively. Furthermore, we computed the fraction of BC mass that can be activated at a certain specific supersaturation ratio based on Kohler Equation.

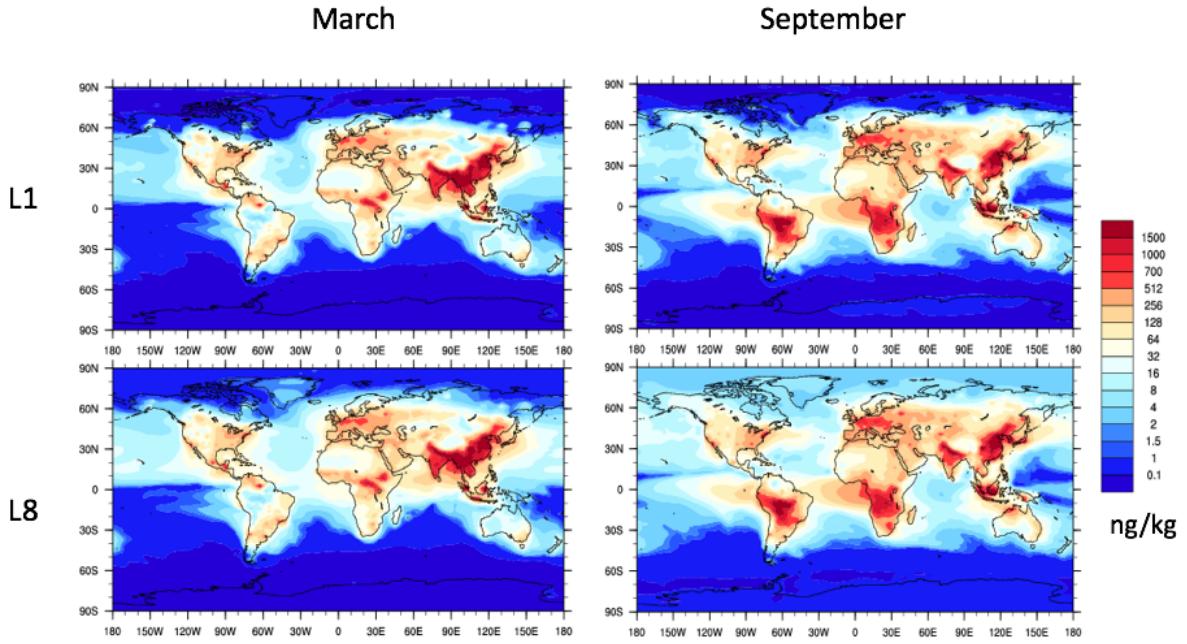


Figure 4.20: BC mass mixing ratio ( $M_{pc}$  and  $M_{accu}$ ) in primary carbon mode (top) and in accumulation mode (bottom), for surface layer, March and September.

In our estimation, we assume that BC cores follow a lognormal size distribution with a fixed standard deviation ( $\sigma=1.4$ ) and a changeable mean diameter ranging from 80 to 150 nm considering the fact that the volume mean size for BC and POM emissions is 134 nm. The number of monolayers ranges from 1 to 23 to cover a wide range of choices. The hygroscopicity value for sulfate is 0.65, and the thickness of one sulfate monolayer is  $4.76 \times 10^{-10}$  m. Figure 4.24 shows the results at supersaturation ratios of 0.3 % and 0.6 % since those values are typical for stratiform cloud. For a 134 nm diameter non-hygroscopic particle, the 8 monolayers of sulfate activates 65.9 % and 99.5% of BC mass with critical supersaturations of 0.3 % and 0.6 % respectively.

## 4.6 Process Analysis of BC Aging

Fierce et al. (2016) has derived a parameterized e-folding time  $\tau_{overall}$  from the approximate

range of condensation growth rate and number concentration for specific locations (Equation 4.1)

$$\tau_{\text{overall}} \approx (k_{\text{cond}} I_{\text{cond}} + k_{\text{coag}} N)^{-1}, \quad (4.1)$$

where  $I_{\text{cond}}$  is the condensation growth rate (in nm/h) and  $N$  is the overall number concentration of aerosol particles (#/m<sup>3</sup>). The condensation coefficient  $k_{\text{cond}} = 0.1$  and the coagulation coefficient  $k_{\text{coag}} = 6 \times 10^{-6}$  cm<sup>3</sup>h<sup>-1</sup> are determined from the regression in Fierce et al. (2016), such that the aging timescale decreases as either  $I_{\text{cond}}$  or  $N$  increase. The above parameterization can be used as a reference for global models. Since condensation and coagulation are independent processes, the overall aging timescale can be separated to two parts, the aging timescale due to condensation:

$$\tau_{\text{cond}} \approx (k_{\text{cond}} I_{\text{cond}})^{-1}, \quad (4.2)$$

and the aging timescale due to coagulation:

$$\tau_{\text{coag}} \approx (k_{\text{coag}} N)^{-1}. \quad (4.3)$$

In our study, we extracted  $I_{\text{cond}}$  and  $N$  from the CAM-chem global model, and computed  $\tau_{\text{overall}}$ ,  $\tau_{\text{cond}}$  and  $\tau_{\text{coag}}$  following Equation 4.1, 4.2, and 4.3, in order to compare them to the modeled simulated aging timescales that are derived from the mass transfer rates.

CAM-chem model simulated aging timescales can be derived by the following equation:

$$\tau = \underbrace{\left( \frac{\frac{\partial m_{\text{BC,cond}}}{\partial t}}{m_{\text{BC,fresh}}} + \frac{\frac{\partial m_{\text{BC,coag}}}{\partial t}}{m_{\text{BC,fresh}}} \right)^{-1}}_{\tau_{\text{cond,CAM}}^{-1}} \quad (4.4)$$

where the mass transfer rate of BC from the primary carbon mode to the accumulation mode

$\frac{\partial m_{BC,cond}}{\partial t}$  (due to condensation) and  $\frac{\partial m_{BC,coag}}{\partial t}$  (due to coagulation) can be extracted from the model, together with the total mass of BC in the primary carbon mode  $m_{BC,fresh}$ . The overall aging timescale  $\tau$ , and analogously, the aging timescale due to condensation  $\tau_{cond,CAM}$  and coagulation  $\tau_{coag,CAM}$  are then estimated according to Equation 4.4.

#### 4.6.1 BC Condensation Growth Rate

In our study, we derived the condensation growth rate  $I_{cond}$  as the volume condensation rate over the total aerosol surface area (in nm/h). As is mentioned in 3.2, the condensation of SOA is reversible, so the total condensation rate can be positive or negative. Since all the fresh BC particles are in the primary carbon mode, we computed the condensation growth rate using the volume condensation rate and surface area extracted from the primary carbon mode. The volume condensation rates are shown in Figure 4.25. A seasonal variation can be observed where the maximum of the condensation rates occurs at 0–30 °S during March, and at 0–30 °N during September. Figure 4.25 also suggests that the distribution of  $I_{cond}$  are quite similar to the distributions of the SOA volume fractions in March, indicating that SOA plays a dominating role in the condensation process in Spring.

#### 4.6.2 CAM-chem v.s. PartMC–MOSAIC Aging Timescales

Figure 4.26 and 4.26 illustrate the CAM-chem modeled v.s. PartMC parameterization derived aging timescales near the surface and in the lower troposphere (857 hPa) respectively. We noted that the aging timescales range from less than one hour (South America) to several days (over the ocean), and these values are broadly consistent with the aging timescales from the PartMC-MOSAIC parameterization near the surface. However, the MAM4 aging timescales are lower than PartMC aging timescales in the lower troposphere as is shown in Figure 4.27. In addition, the aging rates are dominated by the relative abundance of the condensable materials and BC mass, and the condensation of SOA and sulfate plays a

dominating role in BC aging.

We computed the aging timescales in five sensitivity cases using an equivalent of 1, 2, 4, 6 and 8 monolayer-of-sulfate condensation criterion, represented as L1, L2, L4, L6 and L8 respectively (Figure 4.28, 4.29). Judging from the results, when using an 8-monolayer sulfate criterion, the aging timescales near the surface have the closest linear-regression slope to 1 and the highest  $R^2$  value among the five sensitivity cases. At an upper layer (857hPa), however, L6 case have the closest linear-regression slope to 1 and the highest  $R^2$  value, indicating that the CAM-chem aging rates is slow biased.

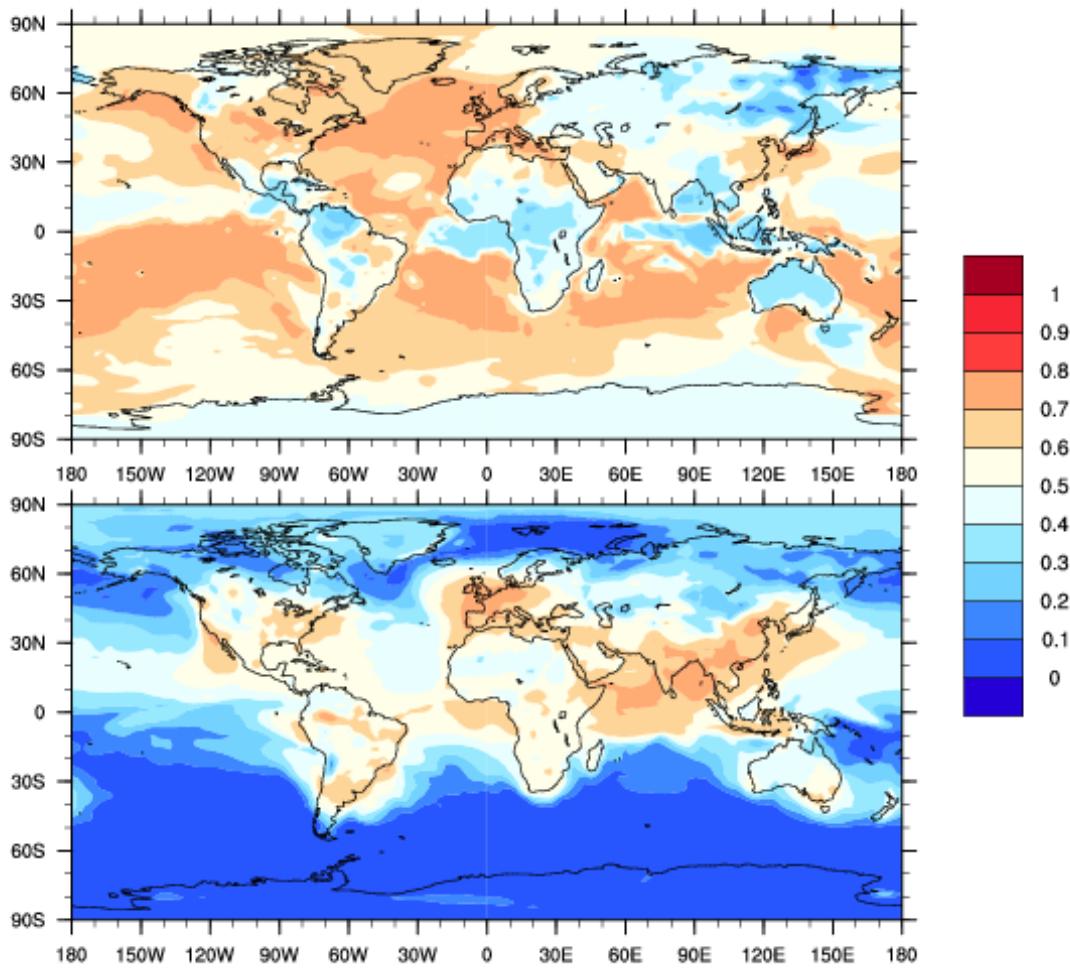


Figure 4.21: BC mass fraction between 90 and 400 nm ( $F_{pc}$  and  $F_{accu}$ ) in the primary carbon mode (top) and accumulation mode (bottom), for surface layer, in March (left) and September (right).

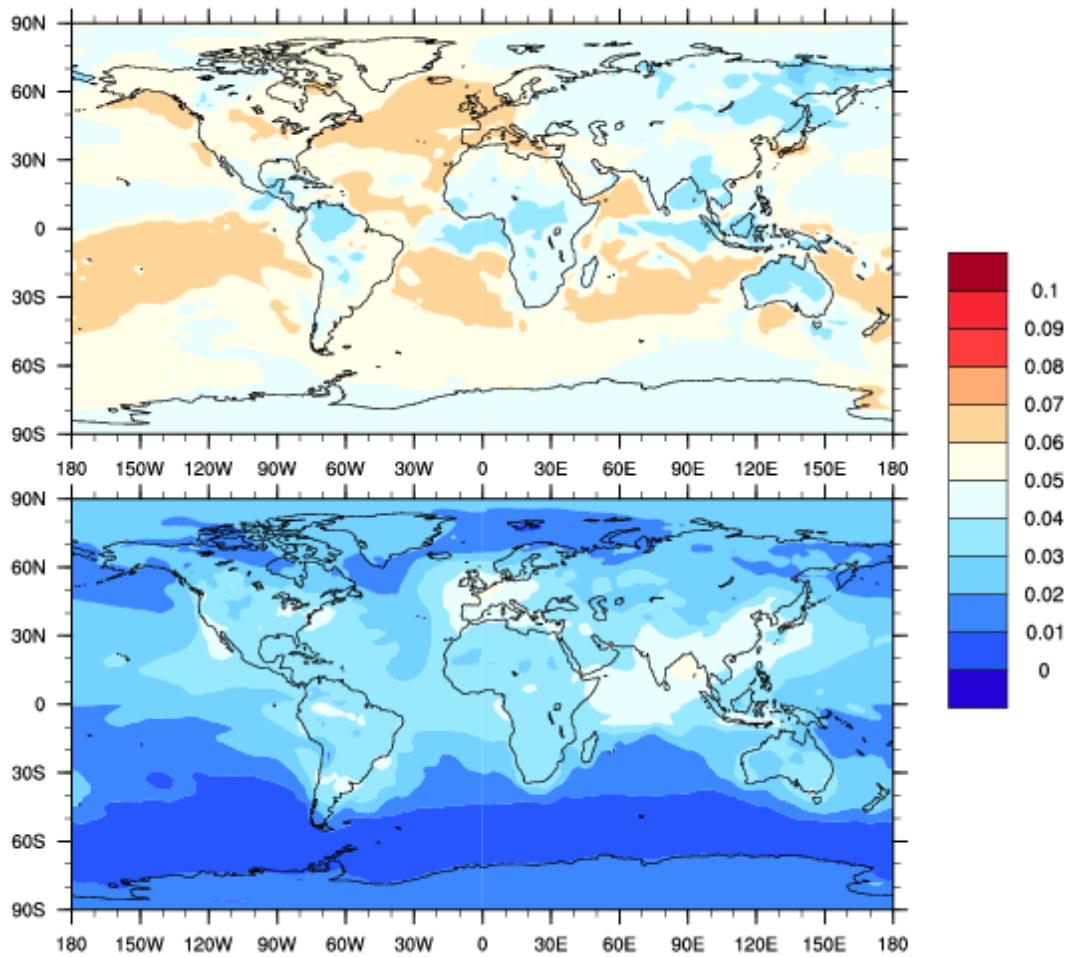


Figure 4.22: Geometric mean diameter of BC core ( $\mu\text{m}$ ) in primary carbon mode (top) and accumulation mode (bottom), for surface layer, in March (left) and September (right).

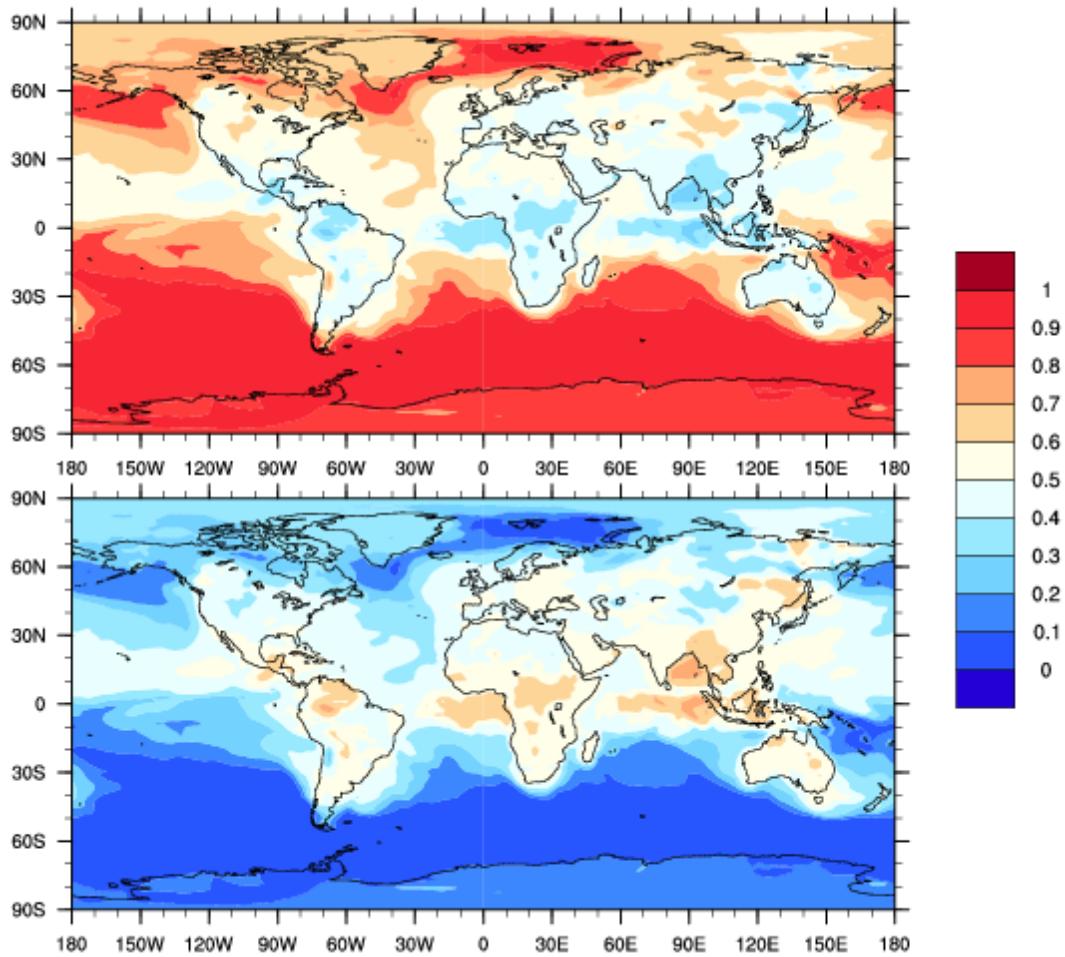


Figure 4.23: Ratio of BC mixing ratio within SP2 size range to total BC mixing ratio within SP2 size range in primary carbon mode  $f_{pc}$  (top) and in accumulation mode  $f_{accu}$  (bottom), for surface layer, in March (left) and September (right).

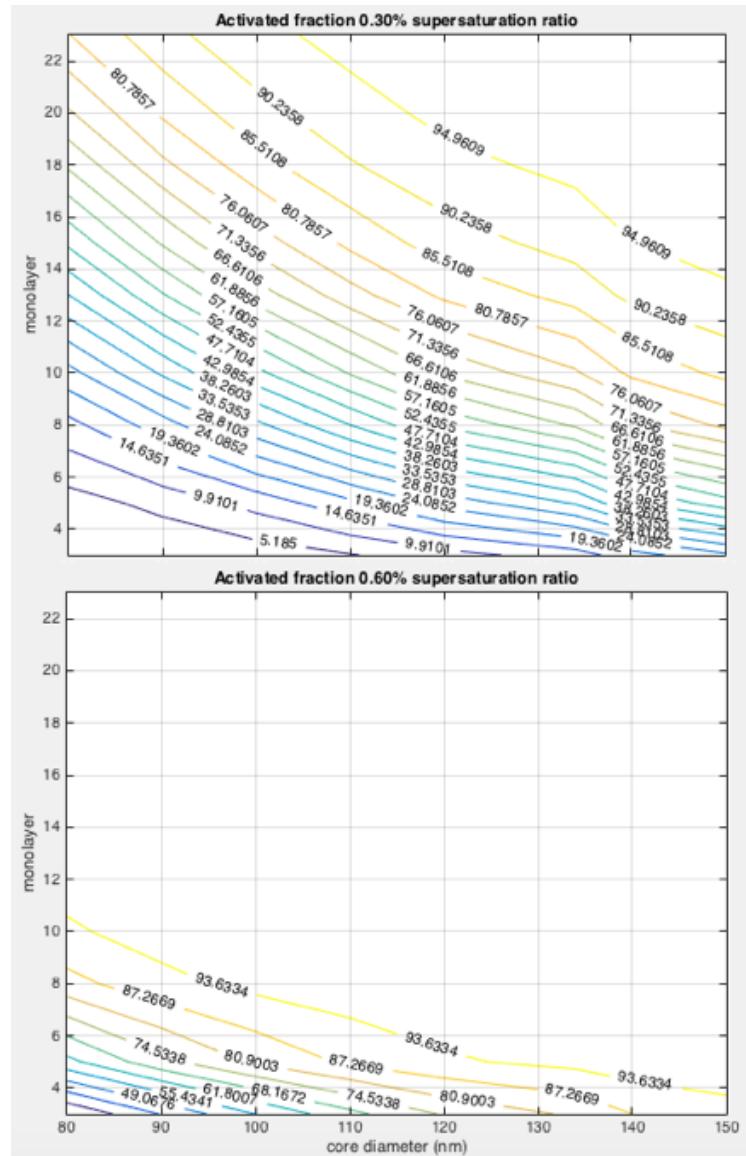


Figure 4.24: Fraction of BC mass that is activated at 0.3% (top) and 0.6% (bottom) supersaturation ratio.

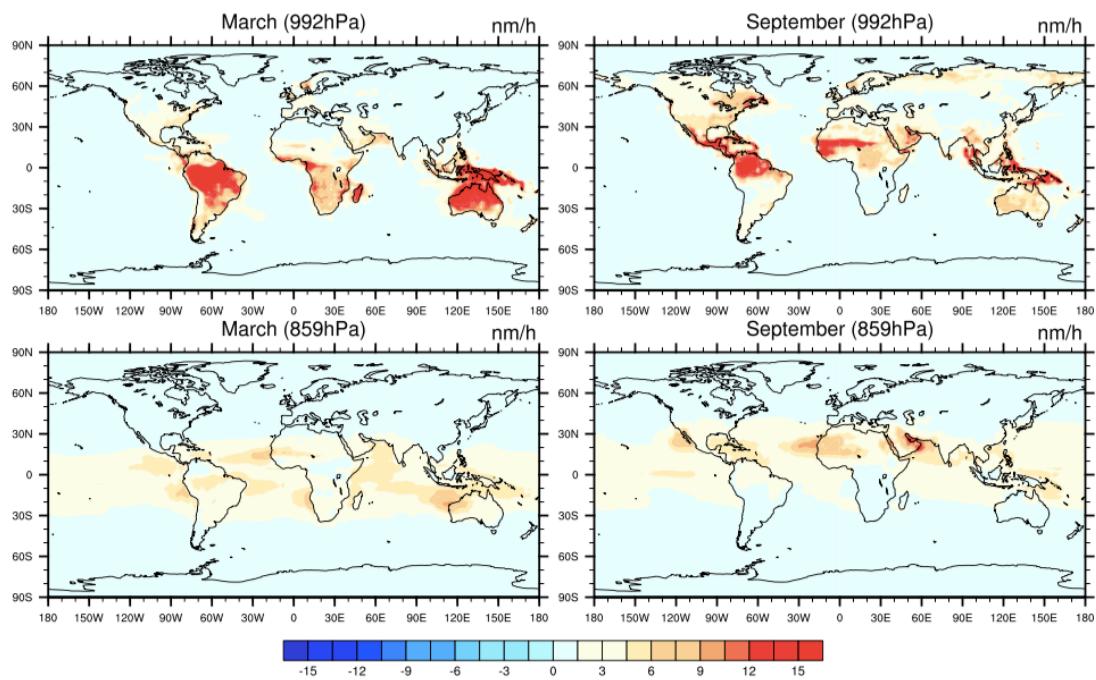


Figure 4.25: Volume condensation rate ( $I_{\text{cond}}$ ) of sulfate and SOA.

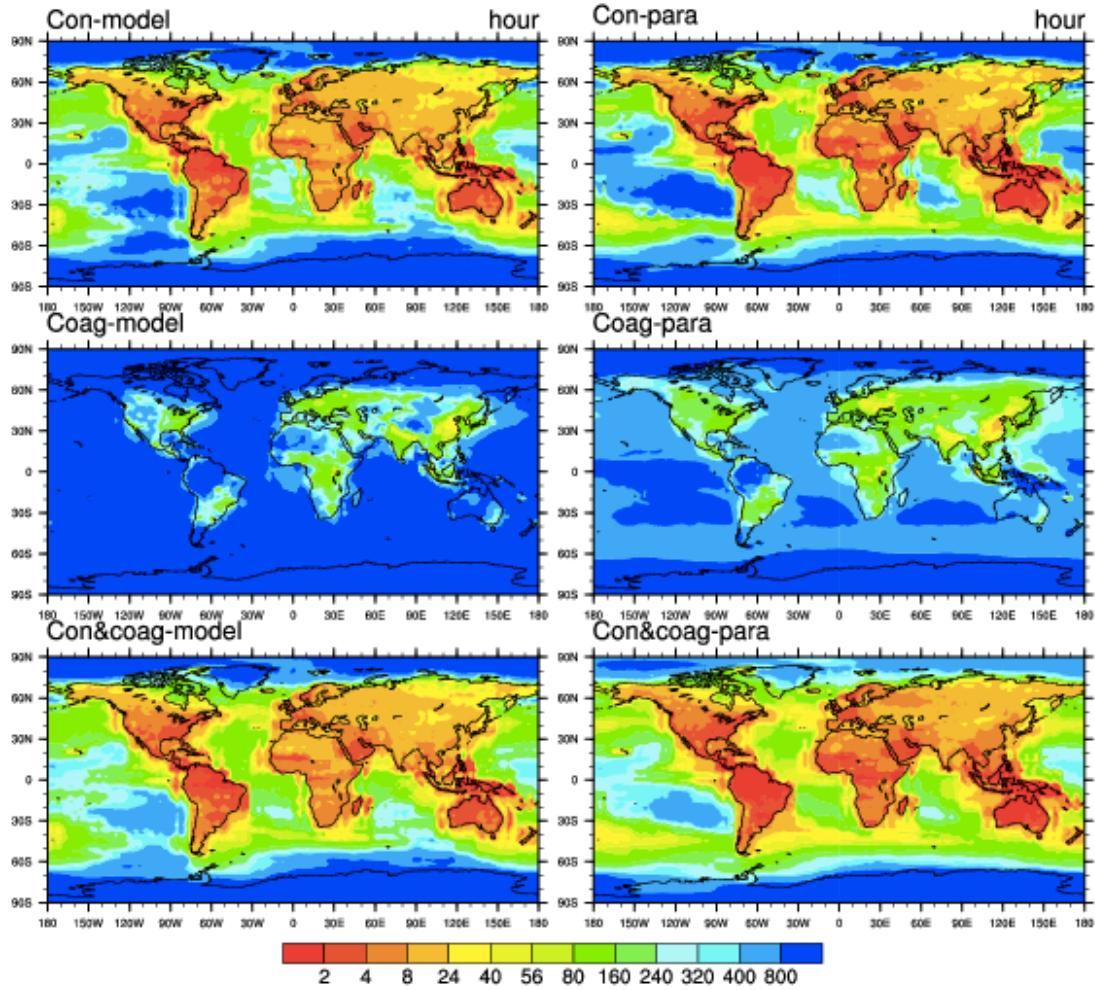


Figure 4.26: Annual BC aging timescales (in hours) near the surface, extracted from the CAM-chem model (left) and computed based on PartMC-MOSAIC parameterization (right). The condensation, coagulation and overall timescales are displayed in the upper panels, middle panels and bottom panels respectively.

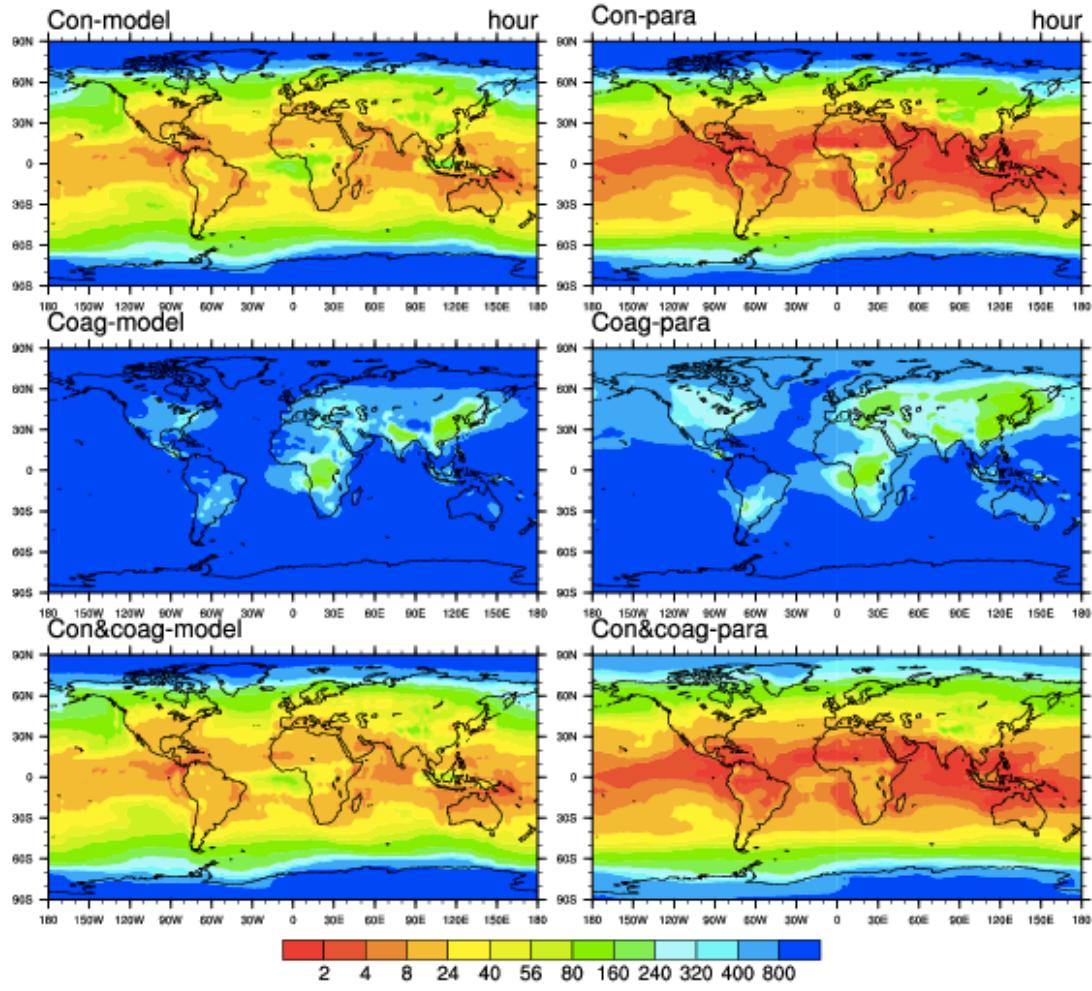


Figure 4.27: Annual BC aging timescales (in hours) at 857 hPa, extracted from the CAM-chem model (left) and computed based on PartMC-MOSAIC parameterization (right). The condensation, coagulation and overall timescales are displayed in the upper panels, middle panels and bottom panels respectively.

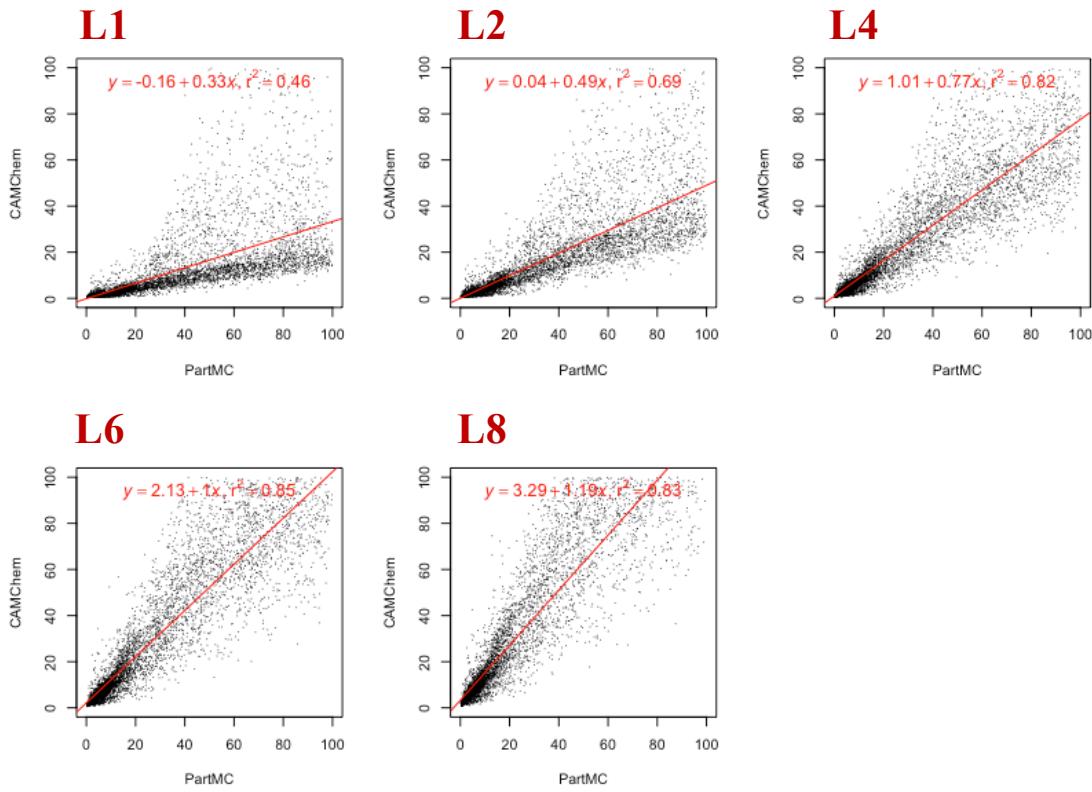


Figure 4.28: Scatter plots of CAM-chem v.s. PartMC–MOSAIC aging timescales (in hour) near the surface in 5 sensitivity cases. L1–L8 represents the number of monolayers (1 to 8) for the condensation criterion.

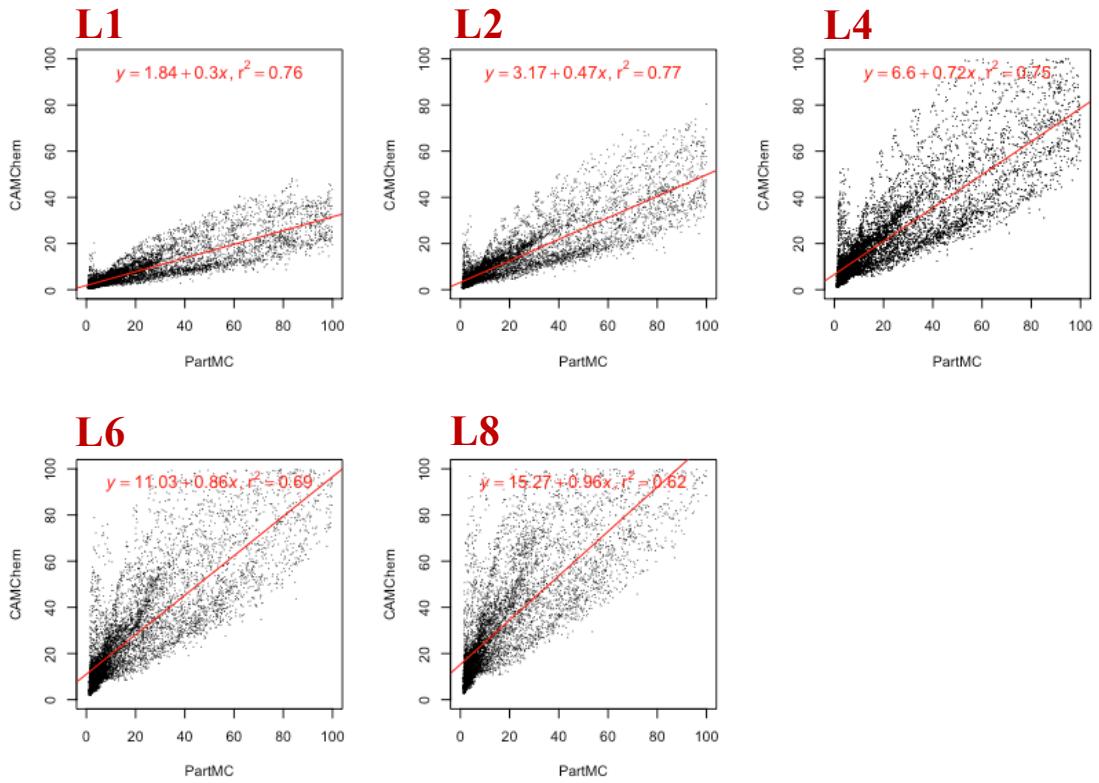


Figure 4.29: Scatter plots of CAM-chem v.s. PartMC–MOSAIC aging timescales (in hour) at 857 hPa in 5 sensitivity cases. L1–L8 represents the number of monolayers (1 to 8) for the condensation criterion.

# Chapter 5

## Summary and Conclusion

In this study, the representation of BC aerosols in CAM–chem MAM4 model has been assessed. Several sensitivity runs were conducted to investigate the extent to which BC burden, mixing states and radiative forcing can be sensitive to the choices of aging parameters. Furthermore, a new method was applied to evaluate the accuracy of BC aging represented in CAM–chem model, by comparing BC aging rates extracted from CAM-chem to aging parameterization based on more detailed particle-resolved simulations with PartMC-MOSAIC. The parameterization were obtained from Fierce et al. (2016).

We used a 4-mode version of the modal aerosol model (MAM4) in CAM–chem, where the aging process was represented by the conversion of BC aerosol particles from the hydrophobic, primary carbon mode to the hydrophilic, accumulation mode after coating with certain amount of species by coagulation and condensation. In order to explore the model sensitivity to the aging criterion, we conducted four simulations from January 1 2010 to December 31 2010 by setting the threshold coating thickness of 1, 2, 4 and 8 of sulfate to determine the mass transfer rate between the two modes. Generally, increasing the number of mono-layers implies that larger amount of sulfate or SOA will be needed to age the primary carbonaceous particles into accumulation mode. Consequently, particles will stay for longer in the primary carbon (no wet deposition) mode before moving into the accumulation mode (subject to wet deposition), hence BC can transport longer distances.

We explored the sensitivity of BC deposition flux and burden to the aging criterion, and computed the relative differences between two cases (L1 v.s. L8). Both wet and dry scavenging rates are most sensitive to the choices of aging criterion at high latitudes, with

the maximum relative differences around 90 % in the Arctic. A seasonal cycle can be observed. Similarly, we found that BC burden is also most sensitive to the aging criterion at high latitudes where the background concentrations are low, with maximum differences in the annually averaged BC mixing ratio of 16% near the surface. For Arctic region, the maximum relative differences can be at high altitudes around 400 hPa. We also analyzed the sensitivity of BC radiative forcing to the aging criterion. The magnitude of annually and spatially averaged radiative forcing is  $0.18 \text{ W m}^{-2}$  in the L1 case, lower than that of  $0.48 \text{ W m}^{-2}$  in the L8 case because less BC are there with larger number of monolayers, and consequently faster aging and wet deposition. The magnitudes are higher in the main source regions (South East Asia, Europe, South America, Africa, etc.) with a maximum of around  $15 \text{ W m}^{-2}$  in both cases.

The mixing state of BC particles with those hydrophilic aerosol components can affect both their CCN/IN activities and radiative properties and should be of our interest. So we computed BC mass fraction and their mixing states, and focused our study in the Arctic region where the climate is more sensitive to BC concentrations. We found that more than 90 % of BC particles are in primary carbon mode (externally mixed) and only less than 10 % of them are in the accumulation mode (internally mixed). We also noticed that SP2 measurements can only catch a portion of BC in both modes because of the detectable size range between 90 and 400 nm. Our results indicated that more than 60 % of BC particles can be detected in March, whereas only less than 20 % of them can be detected in September. This result, albeit preliminary, should be taken into consideration when comparing model simulated BC with SP2 observed data in the Arctic, since the discrepancies may be due to the detective efficiency of SP2 measurements.

Our above results has shown that BC related properties are very sensitive to the choices of aging parameters in the climate model. So it is important to evaluate whether a model representation of BC aging is 'correct'. Currently, limited observed data of aging rates is available, and more generally, it is hard to compare a process to observed data. In CAM-chem, the

treatment of BC aging is based on simplified assumptions, computing mechanic transfer rates by assuming aerosol size distributions, mixing levels and number-of-monolayer criterion. However, PartMC-MOSAIC model tracks the evolution of individual particles (mass constituent aerosol species), where the processes of advection, diffusion and coagulation are modeled stochastically. It has the advantage that no specific aging criterion assumption is required. So in our study, we implemented a new method by exploiting PartMC-MOSAIC parameterizations based on more detailed particle resolved simulations as reference to evaluate the performance of MAM4 aging scheme. We found that CAM-chem annual aging timescales range from less than one hour to several days, where condensation plays a dominating role. The comparison indicates that PartMC-MOSAIC timescales are broadly consistant with CAM-chem timescales near the surface, whereas are lower than CAM-chem aging timescales in an upper layer (857 hPa), indicating faster aging rates. These results can be used as reference for further CAM-chem model development.

This study is mostly based on model simulations. Further work may include comparison between simulated BC burden and mixing states with SP2 data (e.g., HIPPO, FAAM BAe-146, Pallas Global Atmosphere Watch (GAW) ).

# Chapter 6

## Appendix

### 6.1 Configuration of CAM-chem Model

Create a directory for the code, download the code, go into the scripts directory in the source code:

```
1 mkdir ~/cesm1_2_2_CAMChem  
2 cd /glade/u/home/yinruili/cesm1_2_2_CAMChem/script
```

Create a new case:

```
1 setenv CCSMTAG cesm1_2_2_CAMChem  
2 setenv CASE FSTRATMAM4_extract_offmeteo_5  
3 setenv MACH yellowstone  
4 setenv CCSMROOT /glade/u/home/yinruili/${CCSMTAG}  
5 setenv CASEROOT /glade/scratch/yinruili/cases/$CASE  
6 # create a new case using MAM4 and free-running  
7 ./create_newcase -case $CASEROOT -mach $MACH -res f19_f19 -compset FSTRATMAM4  
8 # create a new case using MAM4 with offline-meteorology  
9 ./create_newcase -case $CASEROOT -mach $MACH -res f19_f19 -user_compset GEOS_CAM5%  
    SMA4_CLM40%SP_CICE%PRES_DOCN%DOM_RTM_SGLC_SWAV
```

Edit runtime options using xmlchange: env\_run.xml

```
1 #e.g., start at 2010-01-01, run for 12 months.  
2 ./xmlchange -file env_run.xml -id STOP_N -val '12'  
3 ./xmlchange -file env_run.xml -id STOP_OPTION -val 'nmonth'  
4 ./xmlchange -file env_run.xml -id RUN_STARTDATE -val '2010-01-01'
```

Edit namelist: user\_nl.cam (This file will be created after you set up the case (./cesm\_setup), but it can also be created manually (using cat) before that. I would prefer the latter option).

```

1 #add variables that you would like to see in the output
2 cat <<EOF >! user_nl_cam
3 history_aerosol      = .true.
4 history_amwg         = .true.
5 history_aero_optics = .true.
6 mfilt                 = 1,  1,
7 # nhtfrq = 0, the file will be monthly average
8 # nhtfrq = -24, the frequency is input as 24 hours (daily)
9 nhtfrq                = 0,  -24,
10 # avgflag_pertape = 'A', averaged data
11 # avgflag_pertape = 'I', instantaneous data
12 avgflag_pertape = 'A', 'A',
13 # variables you would like to add into monthly file (fincl1) and hourly file (fincl2)
14 fincl1                ="bcgascon","bcagingcon","bcmasscon","surface","column","bcagingcoag",
15                                "bcmasscoag","numa1","numa2","numa3","numa4", "dgnd_a01", "dgnd_a02", "dgnd_a03",
16                                "dgnd_a04", "dgnw_a01", "dgnw_a02", "dgnw_a03", "dgnw_a04", "dgnumwet1", "dgnumwet2",
17                                "dgnumwet3", "dgnumwet4"
15 fincl2                ="bcgascon","bcagingcon","bcmasscon","surface","column","bcagingcoag",
16                                "bcmasscoag","numa1","numa2","numa3","numa4", "dgnd_a01", "dgnd_a02", "dgnd_a03",
17                                "dgnd_a04", "dgnw_a01", "dgnw_a02", "dgnw_a03", "dgnw_a04", "dgnumwet1", "dgnumwet2",
18                                "dgnumwet3", "dgnumwet4"

```

---

Set offline meteorology:

```

1 inithist = 'MONTHLY'
2 nCDATA      = '/glade/p/cesm/cseg//inputdata/atm/cam/inic/fv/camchem_ic_2008-01-01_1.9
x2.5_L56_c110118.nc'
3 bnd_topo   = '/glade/p/cesm/cseg/inputdata/atm/cam/met/USGS-gtopo30_1.9x2.5
_phys_geos5_c100929.nc'
4 met_data_file = '2010/GEOSS.2_19x2_20100101.nc'
5 met_data_path = '/glade/p/cesmdata/cseg/inputdata/atm/cam/met/GEOSS'
6 met_filenames_list = '/glade/p/cesmdata/cseg/inputdata/atm/cam/met/GEOSS/
GEOSS_filenames_list_c120516.txt'
7 met_max_rlx   = 0.10
8 met_qflx_factor = 0.84

```

---

Configure the model:

```
1 ./cesm_setup
```

---

Modify the source code and add them into ./SourceMode

---

```
1 # For example, I modified two files (al_aero_gasaerexch_sens_icon.F90 and modal_aero_coag
2 .F90), and copied them into the SourceMode directory so that the model can be build
3 with the modified code.
4 cp /glade/u/home/yinruili/cesm1_2_2_CAMChem/scripts/sourcemode/modal_aero_gasaerexch.F90
5 ./SourceMods/src.cam/modal_aero_gasaerexch.F90
6 cp /glade/u/home/yinruili/cesm1_2_2_CAMChem/scripts/sourcemode/modal_aero_coag.F90 ./SourceMods/src.cam/
```

---

Build the model, and then go into the case root and run the model:

```
1 ./CASE.build
2 cd $CASEROOT
3 ./CASE.run
```

---

# References

- Adams, P. J., Seinfeld, J. H., and Koch, D. M. (1999). Global concentrations of tropospheric sulfate, nitrate, and ammonium aerosol simulated in a general circulation model. *Journal of Geophysical Research: Atmospheres*, 104(D11):13791–13823.
- Bauer, S. E., Bausch, A., Nazarenko, L., Tsigaridis, K., Xu, B., Edwards, R., Bisiaux, M., and McConnell, J. (2013). Historical and future black carbon deposition on the three ice caps: Ice core measurements and model simulations from 1850 to 2100. *Journal of Geophysical Research: Atmospheres*, 118(14):7948–7961.
- Baumgardner, D., Kok, G., and Raga, G. (2004). Warming of the arctic lower stratosphere by light absorbing particles. *Geophysical Research Letters*, 31(6).
- Binkowski, F. S. and Shankar, U. (1995). The Regional Particulate Matter Model 1. Model description and preliminary results. *Journal of Geophysical Research*, 100(D12):209.
- Bond, T. C., Doherty, S. J., Fahey, D. W., Forster, P. M., Berntsen, T., Deangelo, B. J., Flanner, M. G., Ghan, S., K??rcher, B., Koch, D., Kinne, S., Kondo, Y., Quinn, P. K., Sarofim, M. C., Schultz, M. G., Schulz, M., Venkataraman, C., Zhang, H., Zhang, S., Bellouin, N., Guttikunda, S. K., Hopke, P. K., Jacobson, M. Z., Kaiser, J. W., Klimont, Z., Lohmann, U., Schwarz, J. P., Shindell, D., Storelvmo, T., Warren, S. G., and Zender, C. S. (2013). Bounding the role of black carbon in the climate system: A scientific assessment. *Journal of Geophysical Research Atmospheres*, 118(11):5380–5552.
- Bond, T. C., Streets, D. G., Yarber, K. F., Nelson, S. M., Woo, J.-H., and Klimont, Z. (2004). A technology-based global inventory of black and organic carbon emissions from combustion. *Journal of Geophysical Research: Atmospheres*, 109(D14).
- Castleman Jr, A., Munkelwitz, H., and Manowitz, B. (1974). Isotopic studies of the sulfur component of the stratospheric aerosol layer 1 2. *Tellus*, 26(1-2):222–234.
- Cheng, C.-H., Lehmann, J., Thies, J. E., Burton, S. D., and Engelhard, M. H. (2006). Oxidation of black carbon by biotic and abiotic processes. *Organic Geochemistry*, 37(11):1477–1488.
- Croft, B., Lohmann, U., and Salzen, K. v. (2005). Black carbon ageing in the canadian centre for climate modelling and analysis atmospheric general circulation model. *Atmospheric chemistry and physics*, 5(7):1931–1949.

- Dentener, F., Kinne, S., Bond, T., Boucher, O., Cofala, J., Generoso, S., Ginoux, P., Gong, S., Hoelzemann, J., Ito, A., et al. (2006). Emissions of primary aerosol and precursor gases in the years 2000 and 1750 prescribed data-sets for aerocom. *Atmospheric Chemistry and Physics*, 6(12):4321–4344.
- Fan, Z., Zhao, Q., Li, T., Yan, J., Ren, Y., Feng, J., and Wei, T. (2012). Easy synthesis of porous graphene nanosheets and their use in supercapacitors. *Carbon*, 50(4):1699–1703.
- Fierce, L., Riemer, N., and Bond, T. C. (2015). Explaining variance in black carbon's aging timescale. *Atmospheric Chemistry and Physics*, 15(6):3173–3191.
- Fierce, L., Riemer, N., and Bond, T. C. (2016). Toward reduced representation of mixing state for simulating aerosol effects on climate. *Bulletin of the American Meteorological Society*, pages BAMS-D-16-0028.1.
- Flanner, M. G., Zender, C. S., Randerson, J. T., and Rasch, P. J. (2007). Present-day climate forcing and response from black carbon in snow. *Journal of Geophysical Research: Atmospheres*, 112(D11).
- Forsström, S., Isaksson, E., Skeie, R., Ström, J., Pedersen, C., Hudson, S., Berntsen, T., Lihavainen, H., Godtliebsen, F., and Gerland, S. (2013). Elemental carbon measurements in european arctic snow packs. *Journal of Geophysical Research: Atmospheres*, 118(24).
- Hakami, A., Henze, D., Seinfeld, J., Chai, T., Tang, Y., Carmichael, G., and Sandu, A. (2005). Adjoint inverse modeling of black carbon during the asian pacific regional aerosol characterization experiment. *Journal of Geophysical Research: Atmospheres*, 110(D14).
- Hamill, P. V., Drizd, T., Johnson, C., Reed, R., and Roche, A. F. (1977). Nchs growth curves for children birth-18 years. united states. *Vital and health statistics. Series 11, Data from the national health survey*, (165):i.
- Highwood, E. J. and Kinnersley, R. P. (2006). When smoke gets in our eyes: The multiple impacts of atmospheric black carbon on climate, air quality and health. *Environment International*, 32(4):560–566.
- Huang, C., Liu, H., Bander, N. H., and Kirby, B. J. (2013). Enrichment of prostate cancer cells from blood cells with a hybrid dielectrophoresis and immunocapture microfluidic system. *Biomedical microdevices*, 15(6):941–948.
- Jacobson, M. Z. (2001). Strong radiative heating due to the mixing state of black carbon in atmospheric aerosols. *Nature*, 409(6821):695–697.
- Kipling, Z., Stier, P., Johnson, C. E., Mann, G. W., Bellouin, N., Bauer, S. E., Bergman, T., Chin, M., Diehl, T., Ghan, S. J., Iversen, T., Kirkev??g, A., Kokkola, H., Liu, X., Luo, G., Van Noije, T., Pringle, K. J., Von Salzen, K., Schulz, M., Seland, y., Skeie, R. B., Takemura, T., Tsagaridis, K., and Zhang, K. (2016). What controls the vertical distribution of aerosol? Relationships between process sensitivity in HadGEM3-UKCA and inter-model variation from AeroCom Phase II. *Atmospheric Chemistry and Physics*, 16(4):2221–2241.

- Koch, D., Schulz, M., Kinne, S., McNaughton, C., Spackman, J., Balkanski, Y., Bauer, S., Berntsen, T., Bond, T. C., Boucher, O., et al. (2009). Evaluation of black carbon estimations in global aerosol models. *Atmospheric Chemistry and Physics*, 9(22):9001–9026.
- Kulmala, M., Laakso, L., Lehtinen, K., Riipinen, I., Maso, M. D., Anttila, T., Kerminen, V.-M., Horrak, U., Vana, M., and Tammet, H. (2004). Initial steps of aerosol growth. *Atmospheric Chemistry and Physics*, 4(11/12):2553–2560.
- Lamarque, J. F., Bond, T. C., Eyring, V., Granier, C., Heil, A., Klimont, Z., Lee, D., Liousse, C., Mieville, A., Owen, B., Schultz, M. G., Shindell, D., Smith, S. J., Stehfest, E., Van Aardenne, J., Cooper, O. R., Kainuma, M., Mahowald, N., McConnell, J. R., Naik, V., Riahi, K., and Van Vuuren, D. P. (2010). Historical (1850–2000) gridded anthropogenic and biomass burning emissions of reactive gases and aerosols: Methodology and application. *Atmospheric Chemistry and Physics*, 10(15):7017–7039.
- Lamarque, J. F., Emmons, L. K., Hess, P. G., Kinnison, D. E., Tilmes, S., Vitt, F., Heald, C. L., Holland, E. A., Lauritzen, P. H., Neu, J., Orlando, J. J., Rasch, P. J., and Tyndall, G. K. (2012). CAM-chem: Description and evaluation of interactive atmospheric chemistry in the Community Earth System Model. *Geoscientific Model Development*, 5(2):369–411.
- Langner, G. (1992). Periodicity coding in the auditory system. *Hearing research*, 60(2):115–142.
- Liousse, C., Penner, J., Chuang, C., Walton, J., Eddleman, H., and Cachier, H. (1996). A global three-dimensional model study of carbonaceous aerosols. *Journal of Geophysical Research: Atmospheres*, 101(D14):19411–19432.
- Liu, D., Whitehead, J., Alfarra, M. R., Reyes-Villegas, E., Spracklen, D. V., Reddington, C. L., Kong, S., Williams, P. I., Ting, Y.-C., Haslett, S., et al. (2017). Black-carbon absorption enhancement in the atmosphere determined by particle mixing state. *Nature Geoscience*, 10(3):184–188.
- Liu, L., Duan, Y., Ma, L., Liu, S., and Yu, Z. (2010). Microwave absorption properties of a wave-absorbing coating employing carbonyl-iron powder and carbon black. *Applied Surface Science*, 257(3):842–846.
- Liu, X., Easter, R. C., Ghan, S. J., Zaveri, R., Rasch, P., Shi, X., Lamarque, J. F., Gettelman, A., Morrison, H., Vitt, F., Conley, A., Park, S., Neale, R., Hannay, C., Ekman, A. M. L., Hess, P., Mahowald, N., Collins, W., Iacono, M. J., Bretherton, C. S., Flanner, M. G., and Mitchell, D. (2012). Toward a minimal representation of aerosols in climate models: Description and evaluation in the Community Atmosphere Model CAM5. *Geoscientific Model Development*, 5(3):709–739.
- Liu, X., Ma, P. L., Wang, H., Tilmes, S., Singh, B., Easter, R. C., Ghan, S. J., and Rasch, P. J. (2016). Description and evaluation of a new four-mode version of the Modal Aerosol Module (MAM4) within version 5.3 of the Community Atmosphere Model. *Geoscientific Model Development*, 9(2):505–522.

- Mauderly, J. L. and Chow, J. C. (2008). Health effects of organic aerosols. *Inhalation Toxicology*, 20(3):257–288. PMID: 18300047.
- McKay, M. D., Beckman, R. J., and Conover, W. J. (1979). Comparison of three methods for selecting values of input variables in the analysis of output from a computer code. *Technometrics*, 21(2):239–245.
- Pacyna, J. M. (1995). *Sources, Particle Size Distribution and Transport of Aerosols*, pages 69–97. Springer Berlin Heidelberg, Berlin, Heidelberg.
- Parungo, F., Nagamoto, C., Zhou, M.-Y., Hansen, A. D., and Harris, J. (1994). Aeolian transport of aerosol black carbon from china to the ocean. *Atmospheric Environment*, 28(20):3251–3260.
- Pöschl, U. (2005). Atmospheric aerosols: composition, transformation, climate and health effects. *Angewandte Chemie International Edition*, 44(46):7520–7540.
- Raatikainen, T., Brus, D., Hyvärinen, A. P., Svensson, J., Asmi, E., and Lihavainen, H. (2015). Black carbon concentrations and mixing state in the Finnish Arctic. *Atmospheric Chemistry and Physics*, 15(17):10057–10070.
- Reddington, C. L., McMeeking, G., Mann, G. W., Coe, H., Frontoso, M. G., Liu, D., Flynn, M., Spracklen, D. V., and Carslaw, K. S. (2013). The mass and number size distributions of black carbon aerosol over Europe. *Atmospheric Chemistry and Physics*, 13(9):4917–4939.
- Riemer, N. (2010). *Introducing semantics*. Cambridge University Press.
- Riemer, N., Vogel, H., and Vogel, B. (2004). Soot aging time scales in polluted regions during day and night. *Atmospheric Chemistry and Physics*, 4(7):1885–1893.
- Riemer, N. and West, M. (2013). Quantifying aerosol mixing state with entropy and diversity measures. *Atmospheric Chemistry and Physics*, 13(22):11423–11439.
- Riemer, N., West, M., Zaveri, R., and Easter, R. (2010). Estimating black carbon aging time-scales with a particle-resolved aerosol model. *Journal of Aerosol Science*, 41(1):143–158.
- Riemer, N., West, M., Zaveri, R. A., and Easter, R. C. (2009). Simulating the evolution of soot mixing state with a particle-resolved aerosol model. *Journal of Geophysical Research: Atmospheres*, 114(D9).
- Schulz, M., Textor, C., Kinne, S., Balkanski, Y., Bauer, S., Berntsen, T., Berglen, T., Boucher, O., Dentener, F., Guibert, S., et al. (2006). Radiative forcing by aerosols as derived from the aerocom present-day and pre-industrial simulations. *Atmospheric Chemistry and Physics*, 6(12):5225–5246.
- Schwarz, J., Spackman, J., Gao, R., Watts, L., Stier, P., Schulz, M., Davis, S., Wofsy, S. C., and Fahey, D. (2010). Global-scale black carbon profiles observed in the remote atmosphere and compared to models. *Geophysical Research Letters*, 37(18).

- Shindell, D., Kuylenstierna, J. C., Vignati, E., van Dingenen, R., Amann, M., Klimont, Z., Anenberg, S. C., Muller, N., Janssens-Maenhout, G., Raes, F., et al. (2012). Simultaneously mitigating near-term climate change and improving human health and food security. *Science*, 335(6065):183–189.
- Tie, X., Brasseur, G., Emmons, L., Horowitz, L., and Kinnison, D. (2001). Effects of aerosols on tropospheric oxidants: A global model study. *Journal of Geophysical Research: Atmospheres*, 106(D19):22931–22964.
- Whitby, E. R. and McMurry, P. H. (1997). Modal Aerosol Dynamics Modeling. *Aerosol Science and Technology*, 27(6):673–688.
- Yue, G. K. and Deepak, A. (1982). Temperature dependence of the formation of sulfate aerosols in the stratosphere. *Journal of Geophysical Research: Oceans*, 87(C4):3128–3134.
- Zaveri, R. A., Easter, R. C., Fast, J. D., and Peters, L. K. (2008). Model for simulating aerosol interactions and chemistry (mosaic). *Journal of Geophysical Research: Atmospheres*, 113(D13).
- Zaveri, T. D., Dolgova, N. V., Chu, B. H., Lee, J., Wong, J., Lele, T. P., Ren, F., and Keselowsky, B. G. (2010). Contributions of surface topography and cytotoxicity to the macrophage response to zinc oxide nanorods. *Biomaterials*, 31(11):2999–3007.
- Zhang, J., Liu, J., Tao, S., and Ban-Weiss, G. A. (2015). Long-range transport of black carbon to the Pacific Ocean and its dependence on aging timescale. *Atmospheric Chemistry and Physics*, 15(20):11521–11535.
- Zuberi, B., Johnson, K. S., Aleks, G. K., Molina, L. T., Molina, M. J., and Laskin, A. (2005). Hydrophilic properties of aged soot. *Geophysical research letters*, 32(1).