

# EESC GR6922: Atmospheric Radiation, Fall 2022

## Homework 3, due November 3

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October 21, 2022

These problems are ambitious and open-ended. My hope is that you'll get a lot out of them. Especially on this assignment you should feel free to collaborate with one another; I ask that you understand any code you use and that your figures and write-ups be your own. As no equations need to be manipulated please type this homework up and turn in a PDF.

As always, please surround your figures and calculations with discussion.

### 1 Creating idealized profiles

Write a small bit of Python to create idealized profiles of temperature, pressure, and water vapor mixing ratio on a discrete grid given a surface temperature. One possible idealization is a troposphere with constant relative humidity following a moist adiabat extending from a surface temperature to  $T = 220\text{K}$ , with an isothermal stratosphere above this. You can make another idealization but should explain your reasoning. Python packages like NCAR's `metpy` may be useful in constructing these profiles.

- On a single panel plot temperature and water vapor mixing as a function of pressure for several values of surface temperature. It'll look more like a sounding if you put pressure (or its logarithm) on the  $y$ -axis.
- On a single panel plot the integrated water vapor path as a function of (air) temperature for several values of surface temperature. It might be more intuitive to have air temperature on the  $y$ -axis, increasing downwards as it would in the atmosphere.

### 2 Line shapes and atmospheric opacity

We've learned how pressure and temperature broadening affect individual absorption lines. This problem illustrates how broadening affects the column-integrated opacity of (simplified) atmospheres.

1. Use `calculate_absxsec_wn()` to compute the spectrally-resolved absorption cross-section ( $\text{m}^2$  per molecule) for carbon dioxide under near-surface and stratospheric conditions. Focus on no more than a few hundred wavenumbers around the  $\nu_2$  vibration-absorption feature at  $667.5\text{ cm}^{-1}$ . What does spectral variation imply about the resolution needed for line-by-line calculations at low pressures vs. high pressures?
2. Use `calculate_absxsec_wn()` to compute the spectrally-resolved absorption cross-section for water vapor under near-surface and stratospheric conditions. It may be most interesting to focus on one spectral region dominated by rotational bands (say around  $500\text{ cm}^{-1}$ ), another in the water vapor window ( $800\text{--}1200\text{ cm}^{-1}$ ), and a third in a spectral region dominated by vibration-rotation lines (say  $1300\text{ cm}^{-1}$ ).
3. Compute the vertically-integrated optical depth due to water vapor and carbon dioxide in your idealized atmosphere. Vertical variations in temperature and pressure will change the spectral absorption via line broadening, and you'll need to convert from cross-sections to total extinction by computing the total number of molecules in each layer. Make plots of both the spectrally-resolved optical depth (a log scale will be helpful) and the transmissivity for a surface temperature representative of the tropics and another of the high latitudes. You may find it useful to double-check against <http://eodg.atm.ox.ac.uk/ATLAS/zenith-absorption>.

### 3 Outgoing longwave radiation is linear in temperature

This problem is inspired by DDB Koll and TW Cronin 2018, “Earth’s outgoing longwave radiation linear due to H<sub>2</sub>O greenhouse effect”, <https://doi.org/10.1073/pnas.1809868115>. Define the longwave (clear-sky) feedback as the change in outgoing longwave radiation with a change in surface temperature

$$\lambda_{LW} = \left. \frac{dF^+}{dT_s} \right|_{TOA} = \frac{dOLR}{dT_s} \quad (1)$$

where  $T_s$  is the surface temperature.

If the range between surface temperature and the (spectrally-resolved) brightness temperature is not enormous, or if temperature variations are roughly constant with height, we might imagine that

$$\lambda_{LW} \approx \frac{d\sigma T_s^4}{dT_s} = 4\sigma T_s^3 \quad (2)$$

In other words, we might expect that  $\lambda_{LW}$  depends fairly strongly on  $T_s$

1. Use `calc_olr_wn()` to compute  $\lambda_{LW}$  at several surface temperatures. Total OLR can be computed by integrating the spectrally-resolved OLR, say with `numpy.trapz()`. The surface temperature of the US Standard Midlatitude summer atmosphere is 294.2 K. Estimate for each surface temperature will require two separate LBL calculations. Compare the values to (2); justify your choice for the temperature at which you evaluate the equation.
2. Plot the spectrally-resolved feedbacks  $\lambda_\nu$  for each pair of temperatures. Why do large regions of the spectrum have such small feedbacks? Which spectral regions have large feedbacks, and why?
3. How and why might feedbacks change at very high surface temperatures in this framework?