## CHEM 304 SPRING 2012 HW/Learning Goals #5: Ozone "Hole" Chemistry & Fundamentals of Tropospheric Chemistry

1. List the non-chemical factors that distinguish Antarctic ozone depletion chemistry from "normal" (i.e. mid-latitude) stratospheric ozone chemistry. Briefly explain their roles.

2. List two reactions that occur on Polar Stratospheric Clouds (PSC's) that convert fairly stable reservoirs into unstable reservoirs. Explain the key difference that makes these "new" reservoirs so "unstable" relative to their "stable" counterparts?

3. As the light returns in the Antarctic spring, do all wavelengths return the same time, or is there a systematic trend? If the latter, describe it and briefly explain its origin. What are the ramifications of a UV deficiency w/ the return of first light? Why does this make the reservoir conversion reactions (in #5) so important?

4. Explain how the "HI ClOx/LOW NOx" situation develops when the first sunlight returns to the south polar stratosphere. Why/how is NOx removed? What reactions occur (with "first light") that liberate ClOx? What specific ClOx species predominates, and how it is formed? What reactions *would* consume it *if* the air was not so "NOx deficient" and/or there was enough deep UV to generate sufficient concentrations of O atoms?

5. List the cycle of reactions that ultimately leads to the (rather sudden) formation of the  $O_3$  "hole" (just after the air is primed for  $O_3$  depletion by the "HI ClOx/LOW NOX" situation, and there is enough near UV (~350 nm) to drive this process).

6. The key reactions in #8 involve (exclusively) members of which catalyst "family"? Are the sources of this family primarily anthropogenic or natural? What are the ramifications: is human activity to blame? Is this a local, regional or global effect?

7. Why must "deep UV" ( $\lambda < 240$  nm) return to the polar stratosphere before significant amounts of O<sub>3</sub> can be regenerated?

8. List the five main (general) processes that may remove or transform any substance released in the troposphere.

9. Natural decay of "biological materials" (i.e. dead plants & animals, etc.), which are composed of large carbon-based molecules, involves breaking up the chains and rings of carbon atoms into small molecules with a single carbon atom. i) If this happens under *aerobic* conditions, what is the main end product of this process? ii) By contrast, what substance is formed under *anaerobic* conditions? iii) Biological waste (e.g. "cow pies") often contains high levels of nitrogen, which results in the release of a simple "reduced" nitrogen substance (even under aerobic conditions – but it is not  $NO_x$ ) – what is it?

10. i) What is the key oxidizing species in the troposphere? ii) Write a sequence two reactions that shows how this species could be produced from any  $O_3$  that diffused into the troposphere from above.

11. NOx enhancement of  $O_3$ : Write a series of three reactions that shows how NO is transformed to NO<sub>2</sub>, ultimately producing  $O_3$ . (The first step requires HO<sub>2</sub>... and also produces OH.)

12. Putting it all together (at least 10 & 11). The diagram below is an attempt to present a holistic, pictorial view of the cyclic processes addressed in #10 and #11. Write chemical equations for each reaction (A-G).

