

Interactive comment on “Deposition of dinitrogen pentoxide, N₂O₅, to the snowpack at high latitudes” by D. M. Huff et al.

Anonymous Referee #2

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This paper presents the first observation based estimates of the N₂O₅ deposition velocity to snowpacks at high latitudes. N₂O₅ is well known to be an important nighttime reservoir of NO_x that can react on surfaces and thus potentially be a sink for NO_x. Surface deposition is thus a likely candidate for N₂O₅ removal, although its importance to the total NO_x budget is difficult to assess given the poor vertical mixing of the nocturnal atmosphere. This issue is especially problematic for high latitudes. Nonetheless, the authors took care in their experimental design, data analysis, and presentation to arrive at a useful data set. I recommend publication in ACP upon their consideration of a few points below.

My general comments concern issues not unknown to the authors, and so I will not belabor them. The gradient method is subject to significant uncertainties, especially

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under stable conditions. The authors attempt to filter the data for periods when the required assumptions are valid, but it appears that leads to the rejection of ~75% of the collected data. Obviously, just because the gradient method cannot be used doesn't mean there wasn't deposition. The authors might wish to discuss the overall representativeness of their measurements in this regard.

I am under the impression that the obtained deposition velocity largely represents a limitation by aerodynamic transfer from the measurement height to the surface or molecular diffusion across some laminar layer, and not a limitation due to actual N₂O₅ loss once in contact with the surface. Is this true, or do measurements of N₂O₅ reactivity on ice surfaces suggest a limitation in this regard?

Other minor comments

Abstract

First sentence is missing "the": "Dinitrogen pentoxide, N₂O₅, is an important intermediate in the nighttime oxidation of NO_x that can react on surfaces."

Line 4-5, Technically, the flux is derived (not observed) from measurements of the N₂O₅ gradient.

Line 10 – 11: ...chemical removal of N₂O₅ that is located within the first few meters?

Introduction Pg 25331, line 15. I think a concluding statement about these different model predictions for the high latitudes might be relevant. Do they predict the same NO_x loss by N₂O₅ in high latitudes as does Dentner and Crutzen?

Pg 25331, line 16. It is interesting that most observations of N₂O₅ atmospheric chemistry have been conducted in mid-latitude regions during summer, when it presumably is least important in a relative sense. Perhaps such an introductory statement to this paragraph would be useful.

Pg 25332, line 16. It doesn't seem possible to know that the deposition was due to

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heterogeneous hydrolysis and not some other reactive loss process.

Methods Pg 25335, line 17: How often was the inlet position switched? Did it spend 15 minutes at one height and then move rapidly to the lower height for 15 minutes? If so, was there an attempt to interpolate both heights to a uniform time base so as to possibly capture some variations that may have occurred over the 30 minute period? If no such attempt was made, perhaps a statement on typical point-to-point differences between two measurements at the same height and how that variation may affect the derived flux would be useful.

Pg 25337, line 10: Introduce Richardson Number index Ri here

Pg 25338, line 9 – 24: I think this discussion should be greatly reduced and streamlined.

Results Pg 25339, line 27: Does the uncertainty quoted here reflect only the standard deviation of the derived values? What was the total number of determinations that went into the average? It seems like you might be underestimating the confidence in the mean value if the quoted range is just the standard deviation.

Discussion

Pg 25340, line 19, I think this first sentence can be cut

It is not clear from the discussion of Apodaca, et al just how big the discrepancy was in required N_2O_5 loss rate in that paper, and whether the derived deposition velocity from this work is really capable of closing that gap quantitatively.

Interactive comment on Atmos. Chem. Phys. Discuss., 10, 25329, 2010.