AMMONIA EMISSION FROM STORED DAIRY MANURE AND ITS LOCAL DEPOSITION

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1. INTRODUCTION

Most of the 151 to 186 Tg of N used in agriculture world-wide end up in other ecosystems (Galloway and Cowling 2002). The transport of NH_3 via the atmosphere is a significant path of this N cycling to non-agricultural systems. Asman and van Jaarsveld (1992) estimated that 44% of the volatilized NH_3 from a source is deposited following an exponential distribution pattern that decreases with distance from the source. Loubet et al. (2006) reviewed the literature on local deposition of NH_3 and concluded that 2 to 60% of the NH_3 from a source is deposited within 2 km.

The objectives of our study was first to report NH_3 emissions over a summer period from dairy cow manure (slurry) stored in a lagoon, secondly to evaluate wind speed and temperature as driving forces of NH_3 emission from this source, and thirdly to document the apparent deposition of NH_3 in the proximity of the lagoon.

2. MATERIALS AND METHODS

Site: A 300-head dairy farm near Lethbridge, Alberta, was used where a manure-storage lagoon was 290 m from the nearest building or trees. The lagoon had a surface area of 41 x 78 m and was surrounded by a berm 2-m in height and was approximately 17-m wide on all sides. The surface of the lagoon typically had a mat of organic material; however, on windy days a portion of the underlying slurry was exposed. The slurry was sampled and the total ammoniacal-N (TAN), NO_x-N, and NO₂-N was evaluated using standard laboratory practices.

Lagoon NH_3 Flux: Wind speed and direction were monitored using a sonic anemometer (CSAT3, Campbell Scientific, Logan, UT). The sonic was positioned 1 m above the surface of the lagoon. The NH₃ mixing ratio (μ mol mol⁻¹) was measured using an open-path laser (GasFinder, Boreal Laser, Edmonton, Alberta) over a path length of 38 m across the eastern section of the lagoon. An infrared thermometer (4000 ALCS, Everst Interscience Inc., Tucsan, AZ) was used to record surface temperature of the lagoon.

All data were filtered and averaged over 15min intervals. These data were used in a dispersion model (WindTrax, Thunder Beach Scientific, Nanaimo, BC) to predict lagoon NH_3 flux.

NH₃ Field Deposition: A second site was set up containing a sonic anemometer and NH₃ laser over an alfalfa field at a distance of 50 m from the lagoon. The sonic anemometer was located at a 2m height above the surface and the laser path was 1.5 m above the surface. Previously, a side-byside comparison of the field and lagoon lasers was made by placing the lasers side-by-side over the lagoon. The regression slope was used to correct the relationship between the two laser concentration values during the study.

The same filtering criteria of the data were applied to the field data as was done for the lagoon data. Only periods where the wind was coming from the direction of the lagoon were considered when determining deposition. The WindTrax model was used to predict NH₃ fluxes from the lagoon and downwind datasets. The difference between the concurrent lagoon and downwind NH₃ fluxes was assumed to be deposition between the lagoon and downwind site.

Statistical Analysis: A path-coefficient analysis was used to determine the impact of wind speed and surface temperature on NH_3 flux (Dewey and Lu 1959).

3. RESULTS AND DICUSSION

The entire data were summarized by constructing a composite 24-h curve using data from all days. On average, the NH_3 flux density over our study

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period was $5.1 \pm 1.6 \text{ g m}^{-2} \text{ d}^{-1}$. The nighttime NH₃ flux density was on average 3.6 g m⁻² d⁻¹ whereas the daytime values were low in the morning and evening, and peaked at midday at 8.6 g m⁻² d⁻¹. These results agree with the NH₃ fluxes from other studies of 4.2 to 6.3 g m⁻² d⁻¹, (Sommer et al. 1993) and 5.3 g m⁻² d⁻¹ (Misselbrook et al. 2005).

The correlation (r) between NH₃ flux and surface temperature was 0.69, while that for wind speed was 0.67. However, surface temperature was also correlated with wind speed (r = 0.25). The path-coefficient analysis indicated the direct influence of surface temperature accounted for 31% of the variability in NH₃ flux, while wind speed was directly responsible for 28% of the variability in NH₃ emissions. Including the indirect effects, surface temperature could explain 45% of the variation in NH₃ emissions, and wind speed 42% of the variation. The tendency is for NH₃ to increase during daylight as surface the temperature and wind speed increase. The importance of temperature as a main factor controlling NH₃ emissions is a result of higher temperature favoring the formation of NH₃ over that of NH₄⁺ in solution; decreasing the solubility of NH₃, and promoting more mineralization of organic matter (Voorburg and Kroodsma 1992). All these changes increase the abundance of gaseous NH₃ at the surface, which increases the surface-to-air NH₃ concentration gradient and therefore the NH₃ emission. The importance of high wind speed is to increase the surface mixing and maintain a surface to air NH₃ concentration gradient that promotes NH₃ emission.

Over the short duration that both the lagoon and downwind laser/sonic units were in place, there were only a few periods that met the criteria of the filtering process and where the wind direction was such that the lagoon was upwind of the laser positioned in the field. In total only 20 values between 19 July and 11 September were available. For these concurrent times, the lagoon NH_3 flux density averaged 5.0 ± 1.1 g m⁻² d⁻¹ while the downwind value was 4.0 \pm 1.0 g m⁻² d⁻¹. These data suggest that about 19% of the emitted NH₃ was captured by the alfalfa crop within 50 m of the lagoon. Close to the lagoon there was an apparent high deposition of NH₃ equivalent to 6.6 kg NH₃-N ha⁻¹ d⁻¹. If this is extrapolated to a longer period of time, it becomes apparent that it would not take many days for the deposited NH₃-N to meet the surrounding crops N requirements for growth.

4. CONCLUSIONS

Quantifying the pathways of NH₃ transfer in agriculture allows evaluation of 'hot-spots' in support of national inventories, and the evaluation of 'net' farm emissions. A key to getting the emissions inventory right is the adoption of accurate emission factors, which should consider net NH₃ emissions (gross emission minus local deposition) from farm. Making the this determination is difficult because the locally deposited NH₃ may be re-emitted as the ambient air NH₃ concentration declines with a change in wind direction, i.e., not associated with a source. Further research is warranted to understand the underlying net effect over time on the long range transport from intensive sources and the immediate area surrounding these sources.

5. REFERENCES

Asman, W. A. H. and H. A. van Jaarsvelt. 1992. A variable-resolution transport model applied for HN_x in Europe. *Atmos. Environ.* 26A:445-464.

Dewey, D. R. and K. H. Lu. 1959. A correlation and path-coefficient analysis of components of crested wheatgrass seed production. *Agron. J.* 51:515-518.

Galloway, J. and E. B. Cowling. 2002. Reactive nitrogen and the world – 20 years of change. *Ambio.* 31:64-71.

Loubet, B., W. A. M. Asman, M. Theobald, O. Hertel, S. Y. Tang, P. Robin, M. Hassouna, U. Dämmgen, S. Genermont, P. Cellier and M. A. Suttton. 2006. Ammonia deposition near hot spots: processes, models and monitoring methods. UNECE Expert Workshop on Ammonia, Edinburgh, 4-6 December 2006. Pp 59.

Misselbrook, T. H., S. K. E. Brookman, K. A. Smith, T. Cumby, A. G. Williams and D. F. McCrory. 2005. Crusting of stored dairy slurry to abate ammonia emissions. *J. Environ. Qual.* 34:411-419.

Sommer, S. G., B. T. Christensen, N. E. Nielsen and J. K. Schjørring. 1993. Ammonia volatilization during storage of cattle and pig slurry: Effect of surface cover. *J. Agric. Sci.* 121:63-71.

Voorburg, J. H. and W. Kroodsma. 1992. Volatile emissions from housing systems for cattle. *Livestock Prod. Sci.* 31:57-70.