## Utilizing the nitrogen isotopic composition of ammonia to investigate regional transport of ammonia emissions: $\delta^{15}$ N-NH<sub>3</sub> values at AMoN sites

J. David Felix<sup>1</sup>, Emily M. Elliott<sup>2</sup>, David Gay<sup>3</sup>

Ammonia (NH<sub>3</sub>) emissions are largely unregulated in the U.S. although wet and dry atmospheric deposition of  $NH_3$  and ammonium  $(NH_4^+)$  can be a substantial source of nitrogen pollution to sensitive terrestrial, aquatic, and marine ecosystems. Despite the adverse effects of excess NH<sub>3</sub> and NH<sub>4</sub><sup>+</sup> deposition (e.g. eutrophication of surface waters, decreased biodiversity, and increased soil acidity), until recently, gaseous NH3 concentrations were not routinely measured as part of the suite of NADP networks. The Ammonia Monitoring Network (AMoN), established in 2007, has rapidly grown to 55 Here, to supplement studies that trace NH3 across local landscapes (e.g. sites. conventionally managed cornfields, confined animal feeding operations, dairy operations), we deployed NH<sub>3</sub> passive samplers at 9 AMoN sites to assess the isotopic composition of NH<sub>3</sub> ( $\delta^{15}$ N-NH<sub>3</sub>) as a regional tracer of NH<sub>3</sub> emission sources. Monthly NH<sub>3</sub> samples from 9 sites were analyzed for nitrogen isotopic composition over a period of a year (7/09 to 6/10). Our results suggest that isotopic compositions of NH<sub>3</sub> at individual AMoN sites generally corresponds with primary regional NH<sub>3</sub> sources. To further explore these spatial patterns, we couple an inventory of the  $\delta^{15}$ N-NH<sub>3</sub> values of NH<sub>3</sub> sources with county-level NH<sub>3</sub> emission inventory (Davidson et al. 2002) to model the average monthly  $\delta^{15}$ N-NH<sub>3</sub> values occurring in U.S. counties. These modeled isotopic compositions are then compared to observed  $\delta^{15}$ N-NH<sub>3</sub> values occurring at individual AMoN sites. This comparison provided insight into possible inaccuracies in the NH<sub>3</sub> inventory and the lack of the modeled isotopic compositions to account for transport of NH<sub>3</sub> sources. These results demonstrate how the nitrogen isotopic composition of NH<sub>3</sub> can be utilized to investigate the source, transport, and fate of NH<sub>3</sub> emissions across varying spatial scales.

<sup>1.</sup> J. David Felix (\*corresponding author), jdf47@pitt.edu, 412 624 8780, Department of Geology and Planetary Science, University of Pittsburgh, 4107 O'Hara St, Pittsburgh, PA 15260

<sup>2.</sup> Emily M. Elliott, <u>eelliott@pitt.edu</u>, Department of Geology and Planetary Science, University of Pittsburgh, 4107 O'Hara St, Pittsburgh, PA 15260

<sup>3.</sup> David Gay, NADP, dgay@illinois.edu 2204 Griffith Drive Champaign, IL 61820-7495

<sup>45</sup>