Effectiveness of Ammonia Reduction on Control of Fine Particle Nitrate

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Abstract. In some regions, reducing aerosol ammonium nitrate (NH_4NO_3) concentrations may substantially improve air quality. This can be accomplished by reductions in precursor emissions, such as nitrogen oxides (NO_x) to lower nitric acid (HNO_3) that partitions to the aerosol, or reductions in ammonia (NH_3) to lower particle pH and keep HNO_3 in the gas phase. Using the
25 ISORROPIA-II thermodynamic aerosol model and detailed observational datasets, we explore the sensitivity of aerosol NH_4NO_3 to gas phase NH_3 and NO_x controls for a number of contrasting locations, including Europe, the US, and China. NO_x control is always effective, whereas the aerosol response to NH_3 control is highly nonlinear and only becomes effective at a thermodynamic "sweet spot". The analysis provides a conceptual framework and fundamental
30 evaluation on the relative value of NO_x versus NH_3 control, and demonstrates the relevance of pH as an air quality parameter. We find that regardless of the locations examined, it is only when ambient particle pH drops below an approximate critical value of 3 (slightly higher in warm and slightly lower in cold seasons) that NH_3 reduction leads to an effective response in $PM_{2.5}$ mass. The required amount of NH_3 reduction to reach the critical pH and efficiently decrease NH_4NO_3

at different sites is assessed. Owing to the linkage between NH_3 emissions and agricultural productivity, substantial NH_3 reduction required in some locations may not be feasible. Finally, controlling NH_3 emissions to increase aerosol acidity and evaporate NH_4NO_3 will have other effects, beyond reduction of $PM_{2.5}$ NH_4NO_3, such as increasing aerosol toxicity and

5   potentially altering the deposition patterns of nitrogen and trace nutrients.

1. Introduction

Global trends of increasing gas-phase ammonia (NH_3) concentrations (Erisman et al., 2008) have multiple environmental implications. As part of the global nitrogen cycle (Fowler et al., 2013), excessive NH_3 deposition promotes alga blooms, degrades water quality, and may be toxic for

5 ecosystems (Krupa, 2003; Camargo and Alonso, 2006). NH_3 is one of the most important atmospheric alkaline species, as it influences the pH of clouds, fogs, precipitation (Wells et al., 1998), and fine particles ($PM_{2.5}$) (Guo et al., 2017c). Agricultural practices, including use of synthetic nitrogen-based fertilizer and domesticated animal manure are the major anthropogenic NH_3 sources (Galloway et al., 2003; Aneja et al., 2009; Zhang et al., 2018). Minor contributions

10 include biomass burning (e.g., forest fires), fossil fuel combustion, and vehicle catalytic converters (Perrino et al., 2002; Behera et al., 2013). Higher temperatures resulting from global warming can also potentially enhance NH_3 emissions (Skjøth and Geels, 2013). Given that fertilizer usage supports food production for about half the global population (Erisman et al., 2008), NH_3 emissions are linked to world population and so expected to increase into the 21th

15 century (Gerland et al., 2014). Compared to the limited regulation of NH_3 emissions, other anthropogenic air pollutants that are linked to acidic atmospheric species, such as sulfur dioxide (SO_2) and nitrogen oxide (NO_x), are regulated through air quality standards, which accounts for their observed decreasing gas and aerosol concentrations in the U.S. (Hand et al., 2012; Russell et al., 2012; Hidy et al., 2014), western Europe, and China (Warner et al., 2017). Decreasing

20 trends of SO_2 and NO_x emissions are expected to continue on global scales throughout the century (IPCC, 2013). The contrast between increasing NH_3 and decreasing SO_2 and NO_x leads to changes in aerosol composition and mass concentration. NH_3 reacts rapidly with the oxidized products of SO_2 and NO_x, sulfuric (H_2SO_4) and nitric (HNO_3) acids, to form ammonium sulfate (($NH_4)_2SO_4$, or other forms such as NH_4HSO_4, $(NH_4)_3H(SO_4)_2$) and ammonium nitrate

25 (NH_4NO_3) aerosols, which globally constitute an important fraction of ambient $PM_{2.5}$ mass (Kanakidou et al., 2005; Sardar et al., 2005; Zhang et al., 2007). These reaction pathways link NH_3 to $PM_{2.5}$ mass and its subsequent impacts on human health (Pope et al., 2004; Lim et al., 2012; Lelieveld et al., 2015; Cohen et al., 2017) and the climate system (Haywood and Boucher, 2000; Bellouin et al., 2011; IPCC, 2013).

A number of studies using regional or global scale models have investigated NH_3 controls as a way to reduce $PM_{2.5}$ mass to meet air quality standards (Erisman and Schaap, 2004; Pinder et al., 2007; Pinder et al., 2008; Paulot and Jacob, 2014; Bauer et al., 2016; Pozzer et al., 2017). The fundamental premise is that reducing NH_3 will increase aerosol acidity (i.e., lower aerosol pH)

5   and prevent the formation of NH_4NO_3, reducing overall $PM_{2.5}$ mass. As a secondary effect, lower pH can also reduce the sulfate production rate, such as the in-cloud SO_2 oxidation by O_3 (Wang et al., 2011; Cheng et al., 2016; Paulot et al., 2017). Use of large-scale models to assess effectiveness of NH_3 controls requires good predictions of a range of pertinent emissions and sinks (NH_3, NO_x, SO_2, and nonvolatile cations), and accurate representation of their applicable

10 atmospheric chemical processes. Thermodynamic modules in different levels of complexity are then applied to determine sensitivities to the precursors (e.g., NH_3, HNO_3). In some cases (Pozzer et al., 2017), the aerosol pH is explicitly determined with an embedded thermodynamic model, such as ISORROPIA-II (Fountoukis and Nenes, 2007). Due to the complexities from all these factors, chemical transport model-predicted responses to changing emissions may not align

15 with observations. For example, the sensitivity of $PM_{2.5}$ pH in the Community Multiscale Air Quality Modeling System (CMAQ) simulations to the mass of crustal material apportioned to the $PM_{2.5}$ size range can have important effects on anticipated responses to these changing emission trends. Vasilakos et al. (2018) have shown that including too much crustal material in $PM_{2.5}$ results in a predicted increasing trend in both aerosol pH and concentrations of NH_4NO_3, which

20 is counter to observations (Weber et al., 2016).

Overall, calculating aerosol pH is a more accurate approach that provides a fundamental understanding of the factors controlling HNO_3-NO_3^- partitioning and therefore enables a direct evaluation of different studies. Furthermore, it is also useful to determine aerosol pH since it has broad application to many other important aerosol processes. For instance, pH is a mediator of

25 many heterogeneous chemical processes, including various acid-catalyzed reactions (Jang et al., 2002; Eddingsaas et al., 2010; Surratt et al., 2010), gas-particle partitioning of species other than HNO_3 and NH_3, such as organic acids and halogens (Fridlind and Jacobson, 2000; Young et al., 2013; Guo et al., 2017a; Nah et al., 2018), and solubility of metals and other nutrient species (Meskhidze et al., 2003; Nenes et al., 2011; Longo et al., 2016; Stockdale et al., 2016; Fang et

30 al., 2017).

In this study, we apply a more direct approach, where measured gas and particle concentrations and the thermodynamic model ISORROPIA-II are used directly in a sensitivity analysis to evaluate the effectiveness of NH_3 emission controls on fine particle mass relative to NO_x control. Contrasts are made between sites that have a wide range in NH_3 concentrations and aerosol composition, with a focus on a one-year dataset collected in Cabauw, the Netherlands (Schlag et al., 2016). This site had year around high NH_3 concentrations (average 7.3 ± 6.0 µg m^{-3}, ~10 ppbv), with nitrate comprising a significant fraction of the fine particle mass (30% NO_3^- of PM_1) and there was a strong seasonal temperature variation. The goal is to establish a transparent and fundamental understanding on when NH_3 emission controls could be an effective way to alter aerosol pH to reduce ammonium nitrate aerosol concentrations, without the use of a full chemical transport model.

2. Methods

2.1 Sampling sites

Cabauw: One-year (July 2012 to June 2013) of online aerosol and gas measurements of inorganic species were made at the Cabauw Experimental Site for Atmospheric Research (CESAR), near the village of Cabauw, the Netherlands. Cabauw is a rural site situated approximately 45 km from the Atlantic Ocean (51.970º N, 4.926º E) and surrounded by agricultural land. Northwestern Europe has fairly high NH_3 concentrations with yearly averages ranging from 1 to 14 µg m^{-3} (median as 4.2 µg m^{-3}) for the Netherlands in 2013, reported by the Measuring Ammonia in Nature (MAN) network (Lolkema et al., 2015). Satellite-derived 14 years average for the western Europe is 3 ppbv (~2.3 µg m^{-3})) (Warner et al., 2017). Cabauw was somewhat higher due to intensive agriculture in the region with observed yearly NH_3 average of 7.3 ± 6.0 µg m^{-3} (~10 ppbv). Site details, instrumentation, and measurement intercomparisons can be found in Schlag et al. (2016). The data used in this analysis is from a monitor for aerosol and gases (MARGA, Applikon Analytical BV) that was operated by the Energy Research Centre of the Netherlands (ECN). The instrument performs online measurements of soluble inorganic gases collected in a continuously wetted-wall denuder, followed by a steam-condensation system for collection of particles. Both the aqueous samples of gases and particles are measured via ion chromatography (Schaap et al., 2011; Rumsey et al., 2014), including NH_3, HNO_3, and HCl, and

particle phase NO_3^-, SO_4^{2-}, Cl^-, NH_4^+, Na^+, K^+, Ca^{2+}, Mg^{2+} alternatively between PM_1 and $PM_{2.5}$, each size sampled hourly (i.e., a two-hour interval for one size; a one-hour interval for gas). Measurement uncertainties were below 10% (Schaap et al., 2011). The detection limits were 0.05, 0.10, 0.08, and 0.01 µg m^{-3} for aerosol ions NH_4^+, NO_3^-, SO_4^{2-}, and Cl^-, respectively, and

5   0.10 and 0.05 µg m^{-3} for the gases HNO_3 and NH_3 (Rumsey et al., 2014). Relative humidity (RH) and temperature (T) data were collected at the 2 m level from the CESAR tower and used to represent ground level meteorological conditions (for an overview see Fig. S7 in Schlag et al. (2016)).

Other Sites: In addition to the Cabauw site, we analyze the effectiveness of NH_3 reduction for a
10   number of contrasting sites where we have already reported on aerosol pH in detail. This includes data from the Southern Oxidant and Aerosol Study (SOAS) (Guo et al., 2015), Wintertime Investigation of Transport, Emissions, and Reactivity (WINTER) (Guo et al., 2016), and California Research at the Nexus of Air Quality and Climate Change (CalNex) study (Guo et al., 2017a). Briefly, the SOAS data was collected at the Southeastern Aerosol Research and
15   Characterization (SEARCH) Centreville ground site, representative of the southeastern US background conditions, from June to July 2013. The WINTER data was sampled from the National Center for Atmospheric Research (NCAR) C-130 aircraft operating from Feb to March 2015 mainly in the northeastern US. The CalNex data was collected from May to June 2010 in Pasadena, California, an urban site that is part of the greater Los Angeles region. As a further
20   contrast for regions of very high NH_3 concentrations, we include an analysis from published data in Beijing during winter haze events in 2015 (Wang et al., 2016), for which pH has also been investigated (Guo et al., 2017c). Table S1 summarizes the conditions at the various sites.

2.2 Thermodynamic modeling

The thermodynamic model ISORROPIA-II (Fountoukis and Nenes, 2007) was used to determine
25   the composition and phase state of an NH_4^+, SO_4^{2-}, NO_3^-, Cl^-, Na^+, Ca^{2+}, K^+, Mg^{2+}, and water inorganic aerosol and its partitioning with corresponding gases. Thermodynamic equilibrium is assumed between fine particles and gases for all semivolatile inorganic species, including particle water and water vapor. Time scales for submicron particles to reach equilibrium are about 30 minutes (Dassios and Pandis, 1999; Cruz et al., 2000; Fountoukis et al., 2009). The
30   model is run in "forward mode" to calculate gas-particle equilibrium concentrations based on the

input of total concentration of inorganic species (e.g., $NH_3 + NH_4^+$, $HNO_3 + NO_3^-$, SO_4^{2-}, Na^+). SO_4^{2-} has no gas pair as it is virtually nonvolatile in the observed temperature ranges of this study (An et al., 2007). The forward mode gives more accurate and robust results than reverse mode since it is much less sensitive to measurement uncertainties (Hennigan et al., 2015).

5   Inorganic ions are also assumed to be only in the aqueous phase (i.e., no solid precipitates). This entails a number of assumptions. First, the ambient RH and the history of the particles exposure to RH result in a deliquesced particle. In many cases, diurnal swings in RH (i.e., the maximum RH in early morning) are generally sufficient to reach the deliquescent point. Furthermore, efflorescence RHs are generally low and rarely reached by the ambient RH (10 to 30%) (Bertram

10   et al., 2011). Thus, a deliquesced particle is often a good assumption when average ambient RH is above 50%. For Cabauw, the one-year mean RH was 81 ± 15 % ($\pm$ SD), with RH reaching up to 90% during diurnal cycles (see Fig. S1a in the supplement) making the presence of liquid phase a reasonable assumption. For the other sites studied, average RHs were all above 55% (Table S1). A second assumption is that most ions are in an aqueous liquid inorganic phase and

15   only minor fractions reside dissolved in a separate liquid organic phase, if it exists. This is supported by very good agreement between observed ammonia gas-particle partitioning with thermodynamic model predictions that do not consider an organic phase. (See Figs. S3 and S4 for this study; similar results are found in other studies (Guo et al., 2015; Guo et al., 2017a; Nah et al., 2018)). Pye et al. (2018) found only minor difference in predicted ammonia partition when

20   an organic phase was considered. It is also assumed that the particles were internally mixed, and that pH did not vary with size. Mixing state of the nonvolatile cations can affect particle composition and pH (Zhu et al., 2016), but the effect on predicted fine particle pH is small if a minor fraction of nonvolatile sulfate is internally mixed with the nonvolatile cations (Guo et al., 2017b), however, it can add uncertainty to predicted nitric acid partitioning (discussed below in

25   Section 2.3). Since there is no data on the mixing state and the mass concentrations (or mole fractions) of nonvolatile cations are generally small (discussed below in Section 3.2 and also see Table S1), internal mixing is assumed in the following analysis.

With increasing pH (e.g., above 2 for oxalate), organic acids can be found at increasing quantities in the particle phase (Nah et al., 2018). However, organic acids are not considered in

30   the ISORRPIA-II pH calculations. In Cabauw, it has been reported that excess NH_4^+ (i.e., NH_4^+ not paired with SO_4^{2-}, NO_3^- and Cl^-) was correlated with (di-)carboxylic organic acids. Excess

NH_4^+ on average constituted only 5% of the NH_4^+ reported by an aerosol mass spectrometer (AMS) so it is likely to have a small effect on predicted pH (Schlag et al., 2017). This is confirmed by the good agreement between measured and ISORROPIA-II predicted NH_3-NH_4^+ partitioning without considering organic acids or other organic species (see section 3.2).

5   Although recent modeling study has suggested that ambient NH_3 concentration can be decreased by as much as 31% in winter and 67% in summer in the US, due to the reactive uptake of NH_3 by secondary carbonyl compounds (Zhu et al., 2018), this process doesn't appear to have an impact on NH_3-NH_4^+ partitioning and predicted pH for the locations in this study. For the winter haze condition in Beijing, the highest pH among the sampling sites, including organic acids (i.e.,

10   oxalate) are reported to decrease pH by at most 0.07, therefore a minor effect (Song et al., 2018).

2.3 NO_x vs NH_3 control to limit $PM_{2.5}$ ammonium nitrate?

Following the various assessments of NH_3 control on $PM_{2.5}$ mass (Erisman and Schaap, 2004; Pinder et al., 2007; Pinder et al., 2008; Paulot and Jacob, 2014; Bauer et al., 2016; Pozzer et al., 2017), we assume the $PM_{2.5}$ inorganic nitrate is mainly in the form of semivolatile ammonium

15   nitrate and negligible in nonvolatile forms, such as $Ca(NO_3)_2$, $NaNO_3$, and similar species, which are generally not found to a large extent in particles smaller than 1 μm. However, it is noted that in locations where concentrations of minerals or sea-salt particle components are high, and the aerosol has aged, formation of semivolatile NH_4NO_3 will be perturbed as the HNO_3 will evolve over time to the more stable largely coarse mode salts (e.g., $CaCl_2$ and NaCl) at the expense of

20   fine mode NH_4NO_3 (see Guo et al. (2017a) for example).

Aerosol organic nitrate species can also contribute to aerosol mass (Farmer et al., 2010; Perring et al., 2013; Xu et al., 2015), and may respond to NO_x control, but are not considered here. For the one-year Cabauw data set analyzed here, 9% of the aerosol nitrate was inferred to be organic nitrate, calculated from the difference in Aerosol Chemical Speciation Monitor (ACSM) nitrate

25   and MARGA-measured nitrate (Schlag et al., 2016). Higher fractions (34% to 44%) have been reported for European submicron aerosols (Kiendler-Scharr et al., 2016). NO_x emission controls could lead to a change in the relative importance of inorganic and organic nitrate (Edwards et al., 2017).

Focusing just on ammonium nitrate, there are two fundamental ways to control $PM_{2.5}$ nitrate;

30   limit the precursors of nitrate aerosol, that is HNO_3, or move the nitrate out of the aerosol by

reducing the aerosol pH (increasing the particle acidity). The equilibrium aerosol nitrate concentration is given by:

$$NO_3^- = \varepsilon(NO_3^-) \times NO_3^T \tag{1}$$

where NO_3^- is the concentration in air of semivolatile aerosol nitrate and $\varepsilon(NO_3^-)$ is the fraction of NO_3^- in the particle phase relative to gas plus particle nitrate ($HNO_3 + NO_3^-$), which is defined

5 as total nitrate, NO_3^T. Eq. (1) is the definition of $\varepsilon(NO_3^-)$. Because $\varepsilon(NO_3^-)$ depends on pH, the premise of NH_3 control is to reduce $\varepsilon(NO_3^-)$ through decreasing particle pH, whereas NO_x emission controls will mainly reduce NO_3^T, although this can also slightly affect pH through aerosol water uptake (discussed below; see Fig. 4 for example).

NO_x Control: Emitted NO_x can undergo a variety of reactions that produce a range of

10 compounds (NO_z), including HNO_3, peroxynitric acid (HO_2NO_2), the nitrate radical (NO_3), nitrous acid ($HONO$), dinitrogen pentoxide (N_2O_5), and both gas (e.g., PAN) and particle phase nitrate and organic nitrate species. Once gas phase HNO_3 or particle phase NO_3^- is formed, equilibrium between the phases will re-establish gas and particle concentrations. HNO_3 is largely formed by NO_2 reaction with the hydroxyl radical (OH), and at night through the nitrate radical-

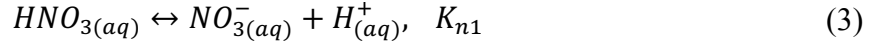
15 N_2O_5 pathway. Modeling studies show that HNO_3 can be the most significant of NO_z species (Atkinson, 2000) and is correlated to NO_x emissions (Shah et al., 2018). Here we assume, to a first approximation, that NO_x mainly produces HNO_3 (either directly through reaction with OH or indirectly through production of N_2O_5) that partitions to the particle to form semivolatile aerosol nitrate and rapidly reaches equilibrium. NO_3^T concentrations are then directly related to

20 NO_x control. Use of more detailed modeling approaches can better assess the relationship between NO_x emissions and NO_3^T. For example, we are not considering competing chemical pathways that lead to organic nitrates, versus inorganic nitrate that is in equilibrium with gas phase HNO_3.

NH_3 Control: The effectiveness of ammonia control in reducing NH_4NO_3 burdens depends on

25 $\varepsilon(NO_3^-)$ and how it varies with pH, actual pH of the ambient aerosol, and the sensitivity of ambient aerosol pH to changes in NH_3 concentration. From thermodynamic equilibrium, $\varepsilon(NO_3^-)$ can be derived from the solubility, reaction (2), and dissociation, reaction (3), of HNO_3:

$$HNO_{3(g)} \leftrightarrow HNO_{3(aq)}, \qquad H_{HNO_3} \tag{2}$$

$$HNO_{3(aq)} \leftrightarrow NO_{3(aq)}^- + H_{(aq)}^+, \quad K_{n1} \tag{3}$$

Assuming the solution is ideal, $\varepsilon(NO_3^-)$ as a function of pH can be predicted solely based on known properties of HNO_3; the HNO_3 Henry's constant, H_{HNO_3}, and the acid dissociation constant, K_{n1} (H_{HNO_3} and K_{n1} are T dependent), ambient T, and particle liquid water content. The latter is often estimated by only considering water associated with inorganic species (W_i; µg m$^-$

5 ^{3}), determined from measured inorganic aerosol components and relative humidity (RH). Liquid water associated with organic species can also be included, but normally have minor influence on pH of much lower hygroscopicity and the logarithmic nature of pH (Guo et al., 2015). A more accurate result may be achieved by using measured particle water concentrations.

By combining the equilibrium of reactions (2) and (3):

$$\varepsilon(NO_3^-) = \frac{H_{HNO_3}^* W_i RT(0.987\times10^{-14})}{\gamma_{NO_3^-}\gamma_{H^+}10^{-pH} + H_{HNO_3}^* W_i RT(0.987\times10^{-14})} \tag{4}$$

10 where 0.987×10^{-14} is a unit conversion factor (from converting atm and µg to SI units), R (J mol^{-1} K^{-1}) is the gas constant and $H_{HNO_3}^* = H_{HNO_3}K_{n1}$ (mol^2 kg^{-2} atm^{-1}) is the combined molality-based equilibrium constant of HNO_3 dissolution and deprotonation, and γ are activity coefficients (equal to 1 if assuming an ideal solution). Derivation of Eq. (4) and references for the temperature dependent equilibrium constants, and similar equations for NH_3 and HCl

15 partitioning, can be found in the supplemental material of Guo et al. (2017a).

3. Results and Discussions

3.1 The nitrate partitioning S Curve

The S curve given by Eq. (4) provides a conceptual basis for the effect of ammonia control, through changes in aerosol pH, on particle nitrate. Fig. 1 shows the characteristic "S-shaped"

20 curve of $\varepsilon(NO_3^-)$ plotted as a function of pH using Eq. (4), for the yearly average conditions in Cabauw and with activity coefficients extracted from ISORROPIA-II ($\gamma_{NO_3^-}\gamma_{H^+} = 0.24$). Including non-ideality shifts the $\varepsilon(NO_3^-)$ S curve to lower pH by approximately 0.6 units (shown as Fig. S2 in the supplement).

Fig. 1 shows that there are 3 pertinent pH regions: 1) low pH, where $\varepsilon(NO_3^-)$ asymptotically approaches 0, and practically all NO_3^T is in the gas phase, 2) $\varepsilon(NO_3^-)$ varies between approximately 0 and 1 and is highly sensitive to pH variations, and, 3) higher pH, where $\varepsilon(NO_3^-)$ approaches 1 and practically all NO_3^T is in the particle phase. This demonstrates that for the one-year average conditions in Cabauw, there is a certain range in ambient pH where NH_3 control to alter ambient pH will result in a change in NO_3^- (i.e., region (2) where pH is between 0 and 3). The greatest change in NO_3^- to a lowering of pH occurs when $\varepsilon(NO_3^-)$ is near 50% (referred to as pH_{50}).

It follows that NH_3 control will only lead to reduction in NO_3^- if ambient particle pH is within region (2) of Fig. 1. If pH is in region (1) there is no need for NH_3 control since pH is sufficiently low that little NO_3^- exists, and if pH is in region (3) the sensitivity of pH to reducing NH_3 will determine the effectiveness of NH_3 controls. For example, NH_3 first needs to be reduced to move particle pH to the transition point between region (2) and (3), where $\varepsilon(NO_3^-)$ starts to drop. (Note that NH_3 control also affects particle mass by changing NH_4^+ concentrations, this is discussed more below in Section 3.4.)

The S curve of Fig. 1 applies for a given situation (see Eq. (4)), which changes as the particle composition or ambient conditions (RH, T) change. For example, if NH_3 concentrations change, the inorganic particle composition changes, which affects particle water and activity coefficients in Eq. (4), resulting in a shift in the $\varepsilon(NO_3^-)$ curve. Thus, these curves provide only a sense of the general state of how NO_3^- responds to changes in NH_3. A full thermodynamic model needs to be run to actually determine the new $\varepsilon(NO_3^-)$ when conditions change. This analysis is provided in the later part of the paper. The S curve, however, provides valuable insight on sensitivity of $\varepsilon(NO_3^-)$ to pH for a given situation (i.e., what region of Fig. 1).

3.2 pH predicted in Cabauw

High concentrations of aerosol inorganic species were observed during the one-year of observations at the CESAR tower. The mass fractions of NO_3^-, SO_4^{2-}, NH_4^+, and Cl^- were on average 30%, 15%, 14%, and 1%, respectively, of the 9.5 µg m^{-3} particle mass (PM_1) (Schlag et al., 2016). The gas-particle partitioning of three semivolatile pairs, NH_3-NH_4^+, HNO_3-NO_3^-, HCl-Cl^-, measured with MARGA are compared with the thermodynamic model predictions (see

section 2 in supplemental material for plots). $PM_{2.5}$ and PM_1 MARGA data sets produce similar results (Fig. S3 versus Fig. S4); here we mainly discuss predictions based on $PM_{2.5}$. Measured and ISORROPIA-predicted partitioning of ammonia was in agreement (NH_3: slope = 1.02, R^2 = 0.997; NH_4^+: slope = 0.97, R^2 = 0.96) (Fig. S3). NO_3^- (slope = 1.01, R^2 = 0.987) and Cl^- (slope = 0.98, R^2 = 0.91) were also in agreement. However, for unknown reasons, gas-phase components of these two species showed significant discrepancies (R^2 of 0.13 to 0.17). We note that it may be associated with the very low gas phase concentrations of these species, in contrast to NH_3. HNO_3-NO_3^- and HCl-Cl^- were dominated by particle phases, $\varepsilon(NO_3^-) = NO_3^-/NO_3^T = 88 \pm 11$ % and $\varepsilon(Cl^-) = Cl^-/(Cl^- + HCl) = 66 \pm 33$ %. The opposite was found for NH_3-NH_4^+, the gas-phase dominated with $\varepsilon(NH_4^+) = NH_4^+/NH_x = 19 \pm 15$ % (total ammonium is referred to $NH_x = NH_3 + NH_4^+$), which is consistent with particle artifacts in the gas collection system possibly affecting HNO_3 and HCl, but less effect on NH_3. Furthermore, a generally better prediction of NH_3-NH_4^+ compared to HNO_3-NO_3^- and HCl-Cl^- partitioning has been observed in our previous studies (Guo et al., 2017a) and is consistent with the lack of a coarse mode sink for NH_3, in contrast to HNO_3 and HCl, which can react with sodium and other nonvolatile cations and bias the equilibrium states between fine particles and gases. In summary, all the semi-volatile inorganic species in the particle-phase (NO_3^-, NH_4^+, and Cl^-) are predicted with high accuracy, as well as NH_3-NH_4^+ partitioning, therefore, particle water and pH predictions by ISORROPIA-II are expected to be reasonable.

As noted above, the presence of water-soluble nonvolatile cations (NVCs, here include Na^+, K^+, Ca^{2+}, Mg^{2+}) can affect the bulk pH analysis. In Cabauw, NVC effects can be assessed by comparing hourly PM_1 and $PM_{2.5}$ data, since these mechanically generated species are largely found in particles larger than 1 µm diameter. Average NVC mole fractions (i.e., NVCs divided by the total inorganic species, not including liquid water), were consistently small, 5.7% for PM_1 and 5.9% for $PM_{2.5}$. However, Na^+ was slightly higher in $PM_{2.5}$ at 0.14 ± 0.25 µg m^{-3}, compared to 0.05 ± 0.09 µg m^{-3} for PM_1. The small and nearly identical fractions of NVCs result in the same predicted pH for PM_1 and $PM_{2.5}$; in both cases pH = 3.7 ± 0.6. Therefore, we focus on the $PM_{2.5}$ in the following discussion due to the similar partitioning predictions and pH for PM_1 and $PM_{2.5}$ (Fig. S3 and S4). A diurnal pattern of ambient particle pH is observed in Cabauw, similar to other studies (Guo et al., 2015). For example, for the nighttime period from 1am to 7am, the

average pH is 3.9, whereas for the daytime period of 1 pm to 6 pm, the pH is ~3.5. The difference is mainly driven by the diurnal variation in liquid water content (see Fig. S1).

3.3 Contrasts in pH and $\varepsilon(NO_3^-)$ between studies

Fig. 2 includes a comparison of $\varepsilon(NO_3^-)$ versus pH for the different locations and seasons (Fig.
5 S5 shows separate plots for each region). The $\varepsilon(NO_3^-)$ curves are plotted based on the campaign average conditions (i.e., T, W_i, and $\gamma_{NO_3}-\gamma_{H+}$; all listed in Table S1). Two sub data sets in Cabauw, summer (June-Aug 2012) and winter (Dec 2012-Feb 2013), are shown together with the one-year whole data set. As seen for Cabauw, lower temperatures (dark blue vs. red vs. orange lines in Fig. 2) shifts HNO_3-NO_3^- partitioning to favor the particle phase due to effect of
10 T on nitric acid Henry's law and dissociation constants, and the explicit effect of T in Eq. 4. For example, at given activity coefficients and liquid water levels, a decrease from 20 °C (~summer) to 0 °C (~winter) shifts $\varepsilon(NO_3^-)$ to lower pH by roughly one unit. The differences between the $\varepsilon(NO_3^-)$ curves are also caused by variations in liquid water, and to a lesser degree by variation in activity coefficients. In general, the summer curves (the right three curves) are at higher pH and
15 the winter curves are at lower pH.

In addition to the S curves, Fig. 2 shows the average ambient particle pH predicted by ISORROPIA-II for each of the studies. Note that pH could also be inferred from the S curve and measured $\varepsilon(NO_3^-)$, but is more uncertain and requires activity coefficients for non-ideality effects. A comparison between Eq. (4)-predicted $\varepsilon(NO_3^-)$ versus pH and observed $\varepsilon(NO_3^-)$ versus
20 ISORROPIA-predicted pH is shown in Fig. S6 and confirms consistency between the ISORROPIA-predicted pH and S curve given by Eq. (4). (A plot of $\varepsilon(NH_4^+)$ vs pH is also shown in Fig. S6). Fine ambient particle pH varies amongst the sites. The pH of 3.7 ± 0.6 in Cabauw is higher than several other regions, such as the SE US (pH = 0.9 ± 0.6), the NE US (0.8 ± 1.0), and the SW US (1.9 ± 0.5), but slightly lower than the China haze ambient particle pH of 4.2. The
25 higher ambient particle pH is generally associated with higher concentrations of NH_3 and NO_3^-. Particle pH is affected by coupling between many variables, hence the need for a thermodynamic model. ISORROPIA-II predicts the overall resulting equilibrium values and associated pH. Particle nitrate has a secondary effect on pH by increasing particle liquid water and diluting H^+ aqueous concentrations, resulting in slightly higher pH. This effect is less pronounced when
30 SO_4^{2-} levels exceed NO_3^-, meaning that liquid water is mainly controlled by nonvolatile SO_4^{2-}.

Thus, NH_3, NO_3^-, and particle pH are coupled. Regions of higher NH_3 will have higher pH which can lead to higher NO_3^- (when in Region (2) of Fig. 1). The highest observed NH_3 (12.8 µg m^{-3}) and NO_3^- (26 µg m^{-3}) concentrations were found for the Beijing haze condition. The Cabauw one-year average NH_3 was lower at 7.3 µg m^{-3}, and NO_3^- was on average of 4.7 µg m^{-3}. The

5 lowest NH_3 and NO_3^- levels were observed in the US studies. For example, 1.37 µg m^{-3} NH_3 and 3.58 µg m^{-3} NO_3^- in the SW US, and only 0.39 µg m^{-3} NH_3 and 0.08 µg m^{-3} NO_3^- in the SE US, both in summer.

The intersection of the $\varepsilon(NO_3^-)$ S curves with ambient particle pH in Fig. 2 (i.e., intersection of vertical line and corresponding site S curve), provide contrast in the average $\varepsilon(NO_3^-)$ at each site,

10 and hence if how much NH_3 control will be needed to shift $\varepsilon(NO_3^-)$ to 50% and corresponding pH of pH_{50}. The lowest $\varepsilon(NO_3^-)$ was found in the SE US at 22% in summer and a higher $\varepsilon(NO_3^-)$ in the NE US in winter at 39%. The Cabauw site also had higher $\varepsilon(NO_3^-)$ in winter (91%) than summer (84%). Additionally, the SW US site observed on average 54% $\varepsilon(NO_3^-)$ in summer and China haze in winter had ~100% $\varepsilon(NO_3^-)$. These data show that in the SE US in summer, $\varepsilon(NO_3^-)$

15 is generally so low that shifting pH by changing NH_3 emissions will not greatly influence NH_4NO_3 concentrations since most is already in the gas phase. Higher NH_3 can increase NH_4NO_3, but large changes in NH_3 are needed in these regions to change pH (Weber et al., 2016). For the SW US summer, NO_3^- partitioning is sensitive to changes in pH with a 54% $\varepsilon(NO_3^-)$. In Beijing winter, substantial decrease in pH is needed to evaporate NH_4NO_3, even

20 more so than Cabauw in winter. For Cabauw, a substantial reduction in ambient pH would be needed to evaporate NO_3^- since the current pH is on the flat zone of the S curve (Region 3), where $\varepsilon(NO_3^-)$ is near 100%. In summer, however, a much smaller reduction in ambient particle pH would result in a decrease in NO_3^-.

3.4 Simulation of particle mass reduction with a thermodynamic model

25 *3.4.1 Sensitivities of pH and nitrate partitioning to NH_3 concentration*

In the above analysis, $\varepsilon(NO_3^-)$ versus pH curves relative to ambient particle pH are used to provide insight on how $\varepsilon(NO_3^-)$ is expected to change with small changes in pH. The S curves are based on the average ambient conditions for each time period, and variables, such as particle water and activity coefficients are held constant. But changes in NH_3 concentration will vary

aerosol composition, liquid water content and the activity coefficients, which in turn modulates the S curve, Eq. (4). To address this, in the following analysis, we run ISORROPIA-II for various input NH_x concentrations, while T, RH, NO_3^T and SO_4^{2-} are held constant, and plot various parameters of interest. This takes into account the various aerosol composition and gas

5   phase species concentrations through considering the partitioning of all semi-volatile species, including water, and how this affects thermodynamic properties, such as activity coefficients.

First, we consider the extent of NH_3 control needed to reduce NH_4NO_3, which depends on the response of pH to changes in ambient NH_3 concentration, which in turn is related to NH_3 emissions (i.e., changes in NH_x). In a previous study, we show that for average conditions at the

10   various sites discussed above, a general rule is that an order of magnitude reduction in NH_3 lowers pH by about one unit (Guo et al., 2017c) ($\Delta pH/\Delta(log_{10}NH_3)$ are listed in Table S1). At the Cabauw site, the responses in pH to changes in NH_3 are similar to these other locations; the linear fitted curves for the semi-log plot in Fig. 3a give slopes of 1.00 in winter, 1.16 in summer and 1.05 for the one-year average (all $R^2 > 0.99$). Fig. 3a also shows predicted pH versus

15   measured NH_3 based on hourly average data. How pH changes with temperature for a constant NH_3 can also be seen in Fig. 3a; higher temperature leads to lower particle pH due to volatilization of semivolatile NH_4^+, NO_3^-, and particle water. The physical explanation for this is that with higher temperature, NH_4^+ is converted to NH_3 and releases one H^+ to the particle phase, whereas NO_3^- is converted to HNO_3 and results in loss of one H^+ from the particle phase. The

20   former process dominates over the latter due to the differences in temperature dependency of equilibrium constants (see Fig. S7) and the greater loss of NH_4^+ from NH_4NO_3 and $(NH_4)_2SO_4$ compared to less loss of NO_3^- only from NH_4NO_3, leading to a net increase in particle H^+ and lower pH. The loss of water associated with NH_4^+ and NO_3^- further reduces pH, as the H^+ becomes more concentrated. The water effect is also seen in the diurnal pH trends (see Fig. S1b).

25   This analysis also permits assessing how $\varepsilon(NO_3^-)$, the sum of NH_4^+ and NO_3^- ($NH_4^+ + NO_3^-$), and $\varepsilon(NH_4^+)$ responds to changes in NH_3. Fig. 3b shows that it takes a factor of 1000 change in NH_3 concentration (~3 pH units) to reduce $\varepsilon(NO_3^-)$ from ~100% to ~0% (i.e. from complete particle-phase to complete gas-phase). Also, a change temperature of ~8 °C shifts $\varepsilon(NO_3^-)$ equivalent to roughly an order of magnitude change in NH_3 concentration. (For reference, ΔT between winter

30   and one-year averages is 6.6 °C and ΔT between one-year average and summer averages is 8.8°C). Fig. 3b and 3c again show that larger reductions in NH_3 are needed in winter compared

to summer to reduce NO_3^-. In Cabauw, only during the highest temperature periods is a NH_3 control policy immediately effective.

Finally, the response of $\varepsilon(NH_4^+)$ to changes in NH_3 is shown in Fig. 3d. The S curves are reversed compared to $\varepsilon(NO_3^-)$ due to opposite base and acid partitioning responses to changes in

5 pH. Thus, lowering NH_3 reduces $\varepsilon(NO_3^-)$, reducing NO_3^- for constant NO_3^T, but raises $\varepsilon(NH_4^+)$ as the particles become more acidic, resulting in relatively more NH_4^+ in the particle phase and less NH_3 in the gas phase. This is important since although we discuss NH_3 emissions, changes in particle pH also affects NH_3 concentrations through changes in gas-particle partitioning, (i.e., $\varepsilon(NH_4^+)$), but it is NH_x that is really changing through emission controls.

10 Finally, Fig. 3d shows that temperature has little effect on the $\varepsilon(NH_4^+)$ versus NH_3 curves. This is because for constant W_i and activity coefficients, the $\varepsilon(NH_4^+)$ versus pH S curves move in the opposite direction with change in temperature than the $\varepsilon(NO_3^-)$ versus pH S curves; $\varepsilon(NH_4^+)$ shifts to a lower pH region and $\varepsilon(NO_3^-)$ shifts to a higher pH region with increasing temperature. This tends to bring the NH_3-NH_4^+ partitioning versus NH_3 curves together and separate the

15 HNO_3-NO_3^- partitioning versus NH_3 curves for different seasons, considering an increase in pH at lower temperature and constant NH_3 shown in Fig. 3a or vice versa.

3.4.2 Effects of NH_3, NO_x, and SO_2 emission control in Cabauw

Here we assess the relative merits of NH_3, NO_x, and SO_2 control on various aspects of $PM_{2.5}$ in Cabauw, again using the full thermodynamic model. Changes in pH, particle water (W_i),

20 $\varepsilon(NO_3^-)$, mass of NH_4^+ plus NO_3^-, and overall $PM_{2.5}$ ion mass are assessed when changes are made to NH_x ($NH_3 + NH_4^+$), NO_3^T ($HNO_3 + NO_3^-$), and SO_4^{2-}, representing control of NH_3, NO_x, and SO_2 emissions, respectively. Each are reduced in steps starting from 0% to a 90% reduction, while holding the other model inputs constant. The results are shown in Fig. 4. The base values are the one-year, summer, and winter average conditions and correspond to 0% reduction in all

25 plots.

The first row in Fig. 4 shows that all parameters respond nonlinearly to NH_x reduction, remaining relatively constant until ~70% NH_x reduction, at which point they start to rapidly decrease. This is a result of the $\varepsilon(NO_3^-)$ versus pH S curve of Fig. 1, where little effect is realized until pH reaches a critical value of about 3 (the horizontal dash line in Figs. 4 pH plots). Once

pH drops below this, the balance between HNO_3 and NO_3^- is sharply shifted towards the gas phase due to the combined effects of reduced particle pH and also reduced particle water (W_i). An approximate 70% reduction in NH_x is required in Cabauw, in winter or based on the yearly average data, to achieve effective reductions in ($NH_4^+ + NO_3^-$) and particle ion mass. In summer,

5 some minor reductions in the mass concentrations occur for small NH_x reductions, since pH is slightly lower in summer (3.3) compared to winter (3.9). Despite the seasonal variations in gas and particle composition, RH and T, all three pH curves (one-year, summer, winter) appear to be similar and show a critical pH of approximate 3; NH_x reduction is more effective for pH below 3 but far less effective for pH above 3, consistent with the simplified analysis above (see Fig. 1).

10 Effects of reducing NO_3^T (the 2nd row, Fig. 4b, i.e., NO_x control) and SO_4^{2-} (the 3rd row, i.e., SO_2 control) show different responses. For NO_x control, holding NH_x and SO_4^{2-} constant, a linear reduction in NO_3^T causes a linear decrease in W_i, ($NH_4^+ + NO_3^-$) and $PM_{2.5}$ ion concentrations simply because $\varepsilon(NO_3^-)$ remains close to 1 so that $NO_3^- \sim NO_3^T$. Then a reduction NO_3^T is just transmitted directly to W_i (SO_4^{2-} is constant so particle hygroscopicity is controlled by NO_3^-),

15 ($NH_4^+ + NO_3^-$) and $PM_{2.5}$ ions. $\varepsilon(NO_3^-)$ is relatively constant (more so in winter) because it is ~100% and so not sensitive to the changes in W_i. Lower W_i does shift the HNO_3-NO_3^- S curve towards a higher pH, but since pH is affected little, and never drops below the critical value of 3, HNO_3-NO_3^- partitioning is barely affected by reducing NO_3^T (i.e., remains in Region (3) in Fig. 1)

20 In the case of SO_4^{2-} reduction, particle pH only increases slightly with substantial SO_4^{2-} reduction due to buffering by NH_3-NH_4^+ partitioning (i.e., NH_4^+ volatility) (Weber et al., 2016; Guo et al., 2017c). ($NH_4^+ + NO_3^-$) decreases slightly due to the loss of associated NH_4^+ due to both the drop in SO_4^{2-} and volatilization caused by reduced particle water. Since SO_4^{2-} is nonvolatile and no gas-particle partitioning is involved, the SO_4^{2-} reduction results in a linear reduction in particle

25 ionic mass, while model input of NH_x and NO_3^T are constant.

Sensitivity test were also performed to investigate the robustness of these results. Considering the observed decreasing trends of SO_2 emissions in many regions (Hand et al., 2012; Hidy et al., 2014; Warner et al., 2017), we tested a cleaner future with less sulfate (20% of the current level, see Fig. S8 in the supplement). Also, since significant changes in global climate and surface land

30 cover can result in a dustier future with more NVCs, we investigated the effect of a 400%

increase in NVCs above the Cabauw levels (see Fig. S9). These two assumed scenarios produce a similar conclusion as the base simulation discussed above, including our finding of a critical pH of 3 and nonlinear response to a NH_x reduction. We do note, however, that in the reduced SO_4^{2-} case, SO_4^{2-} control had nearly no effect on particle ion mass because of the very low SO_4^{2-}

5   concentrations to begin with in the cleaner future scenario.

In summary, the optimal strategy to reduce ammonium nitrate or particle total inorganic ion mass for the current conditions in Cabauw is to control NO_3^T (NO_x emission) since it results in a linear response. Even SO_4^{2-} control is superior over NH_x control to reduce particle ion mass, unless over 70% reduction in NH_x could be achieved. If NH_x is reduced, the effects will be greatest in

10 warmer periods. These are also the times when NH_3 emissions and concentrations are largest regionally or globally (Yamamoto et al., 1988; Warner et al., 2016; Warner et al., 2017; Zhang et al., 2018) (see Table S1 for Cabauw NH_3 levels), and so there may be other benefits to controlling NH_3 emissions at these times, for example, minimizing eutrophication in surface aqueous systems.

15 The above findings in Cabauw are in contrast to results of a global model, which also utilized ISORROPIA-II (Pozzer et al., 2017). They find the impacts of NH_3 emissions on $PM_{2.5}$ mass is strongest in winter for Europe (along with North America, and Asia). Some of the differences are likely attributed to our higher predicted pH in Cabauw of ~3.7 compared to the average pH of Europe predicted in the global model to be near 2 (Pozzer et al., 2017). Thus, we predict

20 conditions above the critical pH of 3, and Pozzer et al. (2017) predicts pH below this value. Difference in pH may be due to meteorological conditions or the concentration of aerosol and gas inorganic species, but it does demonstrate the sensitivity of responses to what the local ambient pH is, and that care should be taken to evaluate predicted particle pH against inferences from ambient measurements. Another thermodynamic model may give a different pH compared

25 to ISORROPIA-II, which may result in a slightly different critical pH (i.e., ~3 in this study). Next, we explore the outcomes of NH_x reductions in other locations and show that NH_3 emission control is more effective in winter than summer.

3.4.3 Effects of NH_3, NO_x, and SO_2 emission control for other locations

NH_x, NO_3^T, and SO_4^{2-} reduction tests were also run for the other sampling sites following the

30 same approach as described above for Cabauw. The model input (period averages) can be found

in Table S1 and the results summarized in Fig. 5. The Cabauw simulations are included in Fig. 5 for direct comparison with the other studies, despite being also plotted in Fig. 4. The average fine particle pH and $\varepsilon(NO_3^-)$ in each study are listed at the top of each plot in Fig. 5 and the plots for the different studies are arranged with increasing ambient pH from left to right. This order is

5 followed in the following discussion.

Fine particles in the eastern US (SOAS and WINTER studies, Fig. 5a and 5b) are the most acidic among the sites, with average pH of approximately 1 due to the lowest NH_3 (and to some minor extent due to small NO_3^-, through its effect on liquid water). In the NE US in winter, NH_x control is most efficient in decreasing $PM_{2.5}$ ion mass since the particle pH and $\varepsilon(NO_3^-)$ (37%) indicate a

10 "sweet spot", where the change in NH_x emission affects NO_3^- immediately. $PM_{2.5}$ ion mass reductions from NO_3^T control and SO_4^{2-} control are similar, since aerosol NO_3^- and SO_4^{2-} are comparable in mass. In the SE US in summer, NO_3^T control is not effective because NO_3^- only contributed 4% to the NH_4^+-SO_4^{2-}-NO_3^- aerosols (Fig. 5a). A small fraction of nitrate aerosol is typically observed in the southeast in summer (Hidy et al., 2014) due to the high temperature and

15 low particle pH. Because of the small NO_3^- fraction and already low pH in summer, NH_x control only leads to minor reductions in particle ionic mass. In contrast, SO_4^{2-} control produces the highest reduction of particle ionic mass since it is the dominant inorganic species (76%) in this region. Therefore, it is more effective to control NH_x in winter in the NE US and SO_4^{2-} in summer in the SE US, a finding consistent with previous studies (Duyzer, 1994; Tsimpidi et al.,

20 2007).

For the southwest US summer (CalNex study, Fig. 5c), since NO_3^- was the most abundant among NH_4^+-SO_4^{2-}-NO_3^- aerosol components, reducing NH_x is the most effective way to reduce $PM_{2.5}$ ion mass as the ambient particle pH is within the range where $\varepsilon(NO_3^-)$ is sensitive to pH. NO_3^T control follows closely in effectiveness, whereas reducing SO_4^{2-} is the least effective. In the

25 WINTER (NE US winter) and CalNex (SW US summer) studies, $PM_{2.5}$ ion mass decreases at a lower rate towards higher levels in NH_x reduction (see Fig. 5b and 5c) due to the nonlinear response in $\varepsilon(NO_3^-)$ to NH_3 concentration (as shown in Fig. 3b or Fig. 2). For instance, when $\varepsilon(NO_3^-)$ drops from 50% to 0%, the sensitivities to NH_3 keeps decreasing asymptotically towards zero. The pH stays nearly flat for the NO_3^T control and SO_4^{2-} control and decreases with NH_x

30 control.

Cabauw winter and Beijing winter haze conditions (see Fig. 5e and 5f) are similar in terms of benefits in reducing particle ionic mass from NH_x, NO_3^T, or SO_4^{2-} controls. This is because of similarities in pH and $\varepsilon(NO_3^-)$ between these sites. For the haze condition in Beijing, NH_x control doesn't produce as much $PM_{2.5}$ ion mass reduction as NO_3^T and SO_4^{2-} controls, unless more than

5   a 60% reduction in NH_x is reached. However, after that PM mass reduction is fast. At 90% NH_x reduction, a decrease of more than half of the particle ionic mass is predicted. NO_3^T and SO_4^{2-} controls produce equivalent results due to the same mass fractions of NO_3^- and SO_4^{2-} (both equal to 36%) and linear response in particle ionic mass. Comparing the pH profiles, the largest reduction in pH is predicted for Beijing haze if reducing NH_x. At 50% NH_x reduction, pH

10   changes from 4.1 to 2.5 in Beijing, whereas pH only changes from 3.9 to 3.3 in Cabauw. This can be explained by differences in $\varepsilon(NH_4^+)$, which is at 60% in Beijing versus 27% in Cabauw.

3.5 Other implications of lowering pH by NH_3 emission control

The benefit of reducing NH_3 emission to reduce ambient $PM_{2.5}$ mass concentrations depends on the conditions at a specific site. While particle pH is lowered during the process, other pH related

15   atmospheric processes are affected. One potentially unintended effect is nitrogen deposition. Nitrogen dry deposition rates depend on particle versus gas phase fractions since there are large differences between gas and particle deposition velocities. For example, the dry deposition velocity of NH_3 is about 1-2 cm s^{-1} over forests, agricultural, or mixed-use land, and 10 times that of NH_4^+ (Duyzer, 1994; Schrader and Brummer, 2014). Also, the dry deposition velocity of

20   HNO_3 is similar to that of NH_3 (Huebert and Robert, 1985). Lowering particle pH through NH_3 reductions will decrease overall reduced nitrogen deposition, but may results in more localized oxidized nitrogen dry deposition, if the lower pH results in NO_3^- evaporation and higher HNO_3 concentrations. Deposition due to wet removal processes are not considered here.

An addition consequence of lowering particle pH is that it can increase aerosol toxicity. Many

25   studies have identified links between strong particle acidity and adverse health endpoints (Koutrakis et al., 1988; Thurston et al., 1994; Raizenne et al., 1996; Gwynn et al., 2000; Lelieveld et al., 2015). We recently showed one way this can happen is due to increased conversion of $PM_{2.5}$ insoluble transition metals to soluble forms by strong acidity (Fang et al., 2017), which increases the particles ability to induce oxidative stress (Ghio et al., 2012).

30   Lowering pH may reduce $PM_{2.5}$ mass but increase overall potential for adverse health effects due

to significantly greater toxicity of soluble metals relative to ammonium nitrate. Finally, lowering pH can also impact the deposition pattern and bioavailability of trace limiting nutrients such as Fe, P, and other metals (Meskhidze et al., 2003; Nenes et al., 2011) with important implications for primary productivity (Meskhidze et al., 2005) and even the oxygen state of the subsurface
5 ocean (Ito et al., 2016).

4. Summary

In this study, we assess the effectiveness of NH_3 control as a way to lower inorganic $PM_{2.5}$ mass based on observational data sets from the US, the Netherlands, and China during different seasons. These sites encompass a diverse range in NH_3 and inorganic aerosol concentrations, and
10 thermodynamic conditions. In all cases, the relative humidities are sufficiently high (average RH > 55%) that a completely deliquesced inorganic phase is a reasonable assumption, which is implicit to the thermodynamic calculations (metastable mode). Focusing on Cabauw, the Netherlands, a site in a region highly impacted by agricultural emissions, we show that the effectiveness of NH_3 control changes with season. In winter, a much larger reduction in NH_3 is
15 required to reduce NO_3^- than in summer, making NO_x control more effective in winter. This is explained by a shift in the HNO_3-NO_3^- partitioning ($\varepsilon(NO_3^-)$) curve to lower pH in winter and pH_{50} (where $\varepsilon(NO_3^-) = 50\%$) further from the actual ambient particle pH. A similar situation is seen in Beijing in winter, where NH_3 emission control would also be less effective. In most other sites investigated, NH_3 control is effective in reducing $PM_{2.5}$ mass, in regions with reasonably
20 high ammonium nitrate concentrations.

The analysis presented here provides a conceptual and direct evaluation of how the inorganic gas-particle system can be expected to respond to changes in NH_3 emissions, and how it contrasts to NO_x control. The approach relies on the single HNO_3-NO_3^- partitioning equation and the use of a thermodynamic model to predict pH. Other approaches are also often used to address
25 this question. Chemical transport models with imbedded thermodynamic sub-modules (such as ISORROPIA) can provide a more detailed analysis that includes other possible impacts of the emission controls, such as ammonia and nitrate deposition and associated environmental impacts. However, the various uncertainties associated with the many simulated processes involved in these models (e.g., emissions and processing) can affect the predicted results and

obscure the fundamental partitioning processes. With the more transparent and accessible approach presented here, this is less an issue. Both approaches have benefits, but whichever analysis is utilized, it is always useful to explicitly report estimated particle pH as it allows assessment of the predictions and provides contrasts between studies at specific sites.

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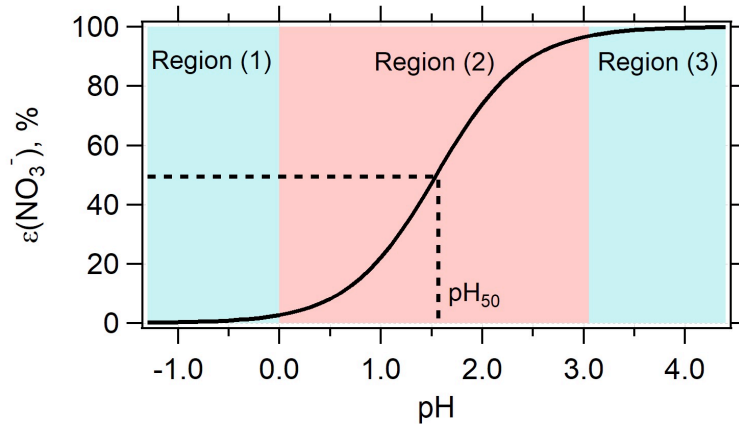
Figure 1. Predicted particle phase fraction of total nitrate, $\varepsilon(NO_3^-)$, versus pH for one-year

5 average condition in Cabauw based on Eq. (4). The blue-color zone denotes where HNO_3-NO_3^- (nitric acid-nitrate) partitioning is not affected by changes in pH, while the red-color zone shows the region where adjusting pH will change HNO_3-NO_3^- partitioning, hence NO_3^- concentration. Greatest sensitivity NO_3^- occurs at $\varepsilon(NO_3^-) = 50\%$, corresponding to pH_{50}.
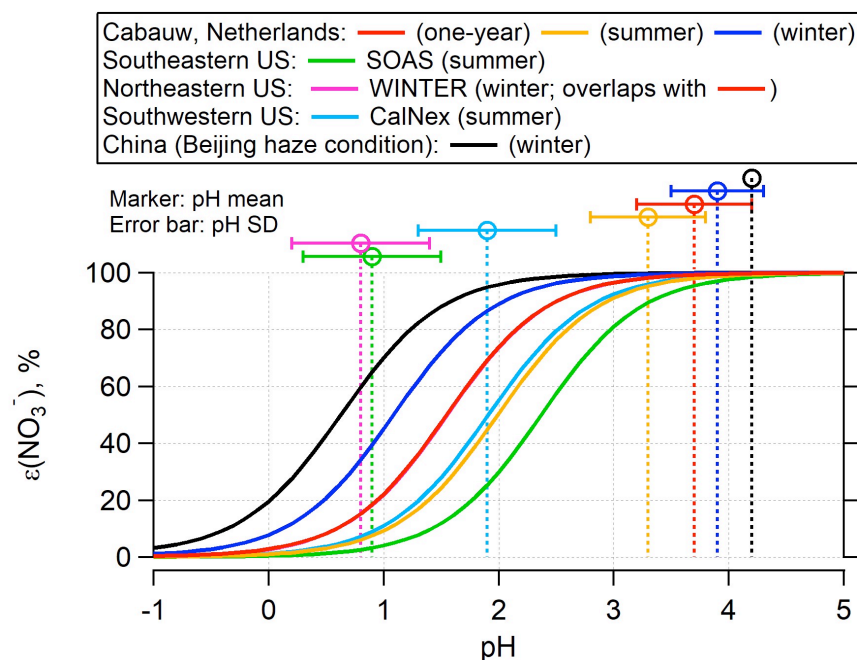
Figure 2. ε(NO$_3^-$) versus pH for various field studies based on the average temperature, liquid water, and activity coefficients for each study, according to Eq. (4). The WINTER study curve overlaps completely with the Cabauw one-year average curve in red color. The input can be found in Table S1. Vertical lines are the study average ambient fine particle pH calculated with ISORROPIA-II and error bars show the variability in pH as one standard deviation. S-curves and ambient pH for each site or season can be matched by color. For a more direct comparison between seasons at a specific region, supplemental Fig. S5 shows separate curves and ambient pH plots.
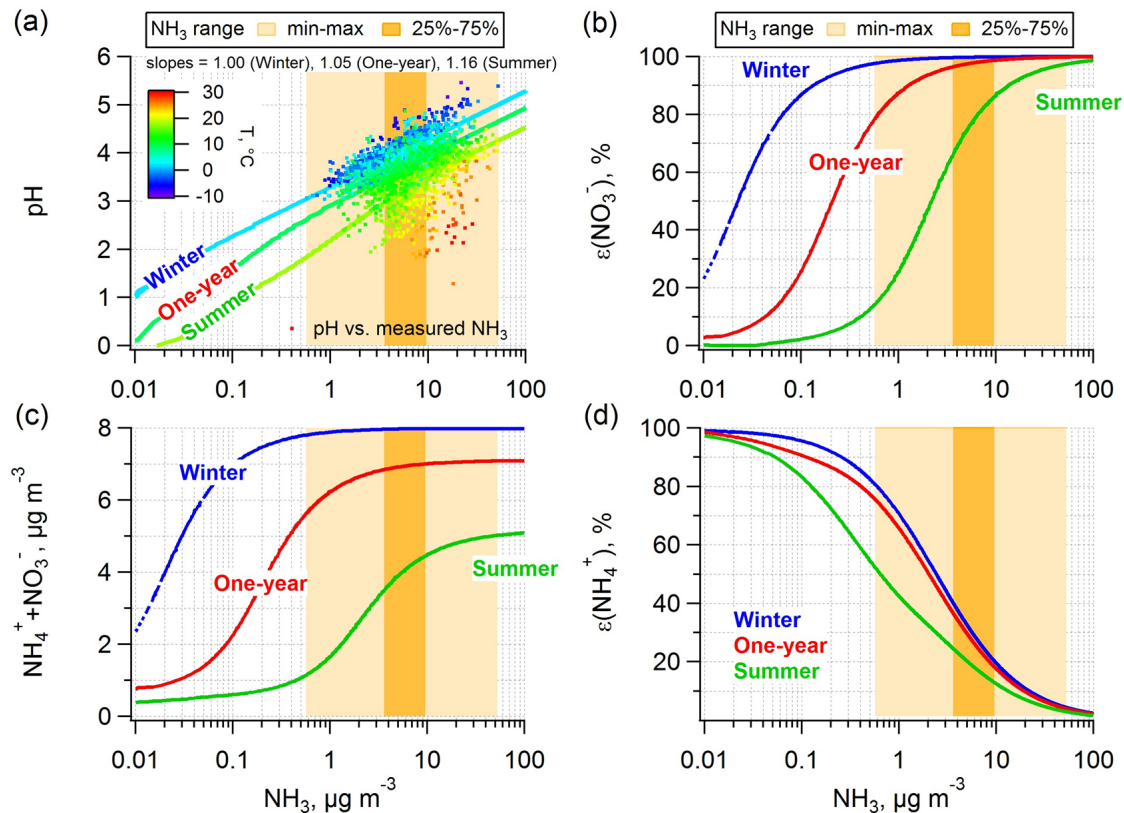
Figure 3. Prediction of (a) particle pH, (b) particle phase fractions of total nitrate, $\varepsilon(NO_3^-)$, (c) ammonium and nitrate mass concentration, (d) particle phase fractions of total ammonium, $\varepsilon(NH_4^+)$ for a wide range of ammonia. The simulations are based on the one-year (July 2012-June 2013), summer (June-Aug 2012), and winter (Dec 2012-Feb 2013) average conditions at the Cabauw site with NH_x ($NH_4^+ + NH_3$) left as a free variable. The measured NH_3 ranges for the one-year span are also shown as the lighter (min-max) and darker (25%-75% percentiles) orange-color zones. Plot (a) also includes the predicted pH versus measured NH_3 data for the entire study and colored by ambient temperature.
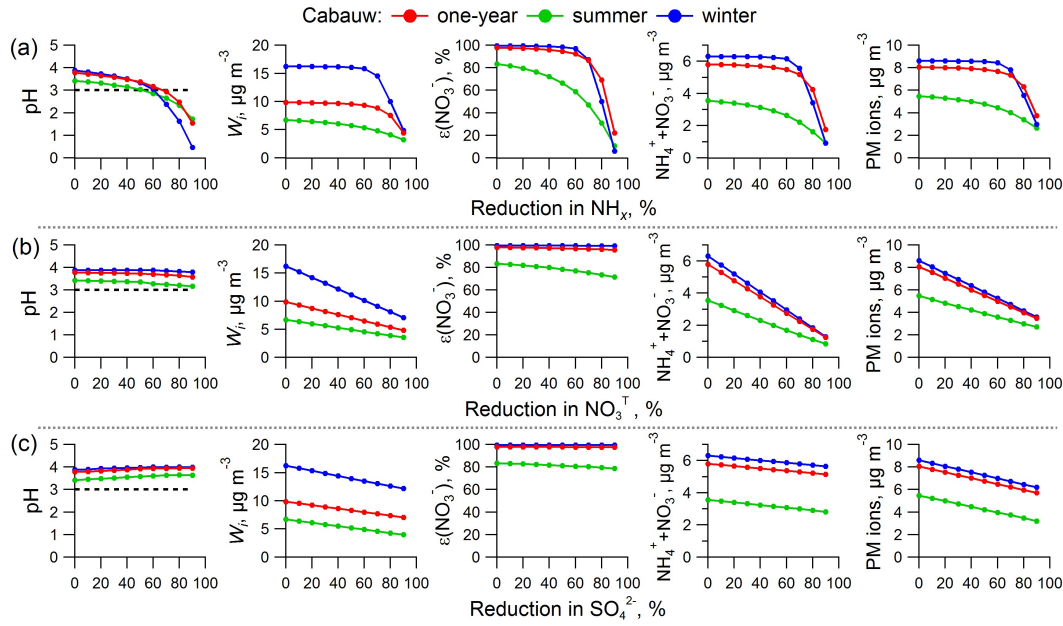
Figure 4. ISORROPIA-predicted $PM_{2.5}$ pH (1st column), liquid water content (W_i, 2nd column), $\epsilon(NO_3^-)$ (3rd column), ammonium and nitrate (4th column), and aerosol inorganic mass concentrations (5th column) as a function of changes in NH_x ($NH_4^+ + NH_3$, 1st row), NO_3^T ($NO_3^- + HNO_3$, 2nd row), and SO_4^{2-} (3rd row). Simulations are based on average conditions of one-year, summer, and winter observational data in Cabauw, the Netherlands, and changing only NH_x, NO_3^T and SO_4^{2-} from the average conditions. The black dash lines in the pH figures identifies the critical pH value of 3.
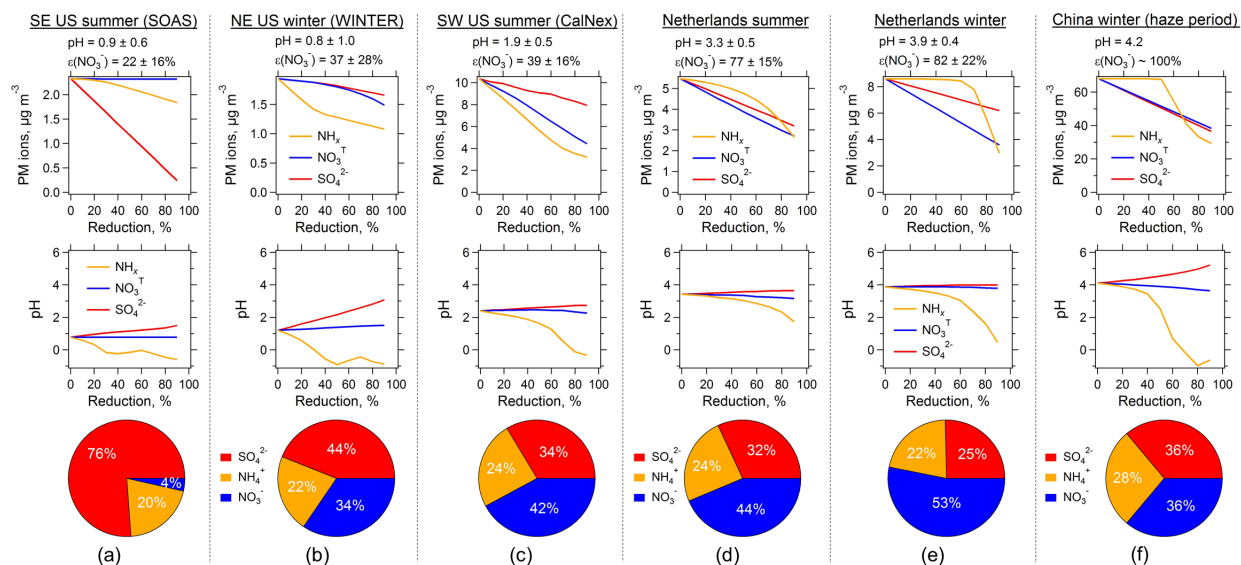
Figure 5. Response of predicted PM$_{2.5}$ inorganic mass concentration (1st row) and pH (2nd row) to reduced levels of NH$_x$ (NH$_3$ + NH$_4^+$), NO$_3^T$ (HNO$_3$ + NO$_3^-$), and SO$_4^{2-}$ for several studies including: (a) the southeastern US summer at a rural ground site in Centreville, AL (SOAS study), (b) the northeastern US during winter (WINTER aircraft study), (c) the southwestern US summer at an urban site in Pasadena, CA (CalNex study), (d) & (e) the Netherlands summer and winter conditions at an rural site in Cabauw from this study, and (f) polluted winter conditions (haze) in Beijing, China. For each case, the average fine ambient particle pH and ε(NO$_3^-$), prior to the reductions, are shown above the figures, with the columns ordered with increasing ambient particle pH from left to right. PM$_{2.5}$ mass fractions of NH$_4^+$-SO$_4^{2-}$-NO$_3^-$ based on study averages are shown as pie graphs along the bottom.