

High efficiency of livestock ammonia emission controls in alleviating particulate nitrate during a severe winter haze episode in northern China

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Abstract. Although nitrogen oxide (NO_x) emission controls have been implemented for several years, northern China is still facing high particulate nitrate (NO_3^-) pollution during severe haze events in winter. In this study, the thermodynamic equilibrium model (ISORROPIA-II) and the Weather Research and Forecast model coupled with chemistry (WRF-Chem) were used to study the efficiency of NH₃ emission controls on alleviating particulate NO₃⁻ during a severe winter haze episode. We found that particulate- NO_3^- formation is almost NH₃-limited in extremely high pollution but HNO₃limited on the other days. The improvements in manure management of livestock husbandry could reduce 40 % of total NH₃ emissions (currently 100 kt month⁻¹) in northern China in winter. Consequently, particulate NO₃⁻ was reduced by approximately 40 % (on average from 40.8 to $25.7 \,\mu g \,m^{-3}$). Our results indicate that reducing livestock NH₃ emissions would be highly effective in reducing particulate NO_3^- during severe winter haze events.

1 Introduction

In northern China (including Beijing, Tianjin, Hebei, Shandong, Shanxi and Henan), severe haze pollution events occur frequently during wintertime, with the concentration of $PM_{2.5}$ (particles with an aerodynamic diameter less than 2.5 µm) reaching hundreds of micrograms per cubic meter and SIA (secondary inorganic aerosol) accounting for more than 50% of $PM_{2.5}$ (Zheng et al., 2016; Tan et al., 2018). To mitigate fine-particle pollution, the Chinese government has been taking strong measures to control SO₂ emissions (http: //www.gov.cn/zwgk/2011-12/20/content_2024895.htm, last access: 5 January 2019). Since 2007, SO₂ emissions have been reduced by 75% in China (Li et al., 2017). Consequently, the particulate-sulfate concentration has also been declining continuously in the past decade (Geng et al., 2017).

Although NO_x emissions in 48 Chinese cities decreased by 21% from 2011 to 2015 (Liu et al., 2017a), unfortunately, no obvious decreasing trend for particulate NO₃⁻ has been observed in northern China during recent years (Zhang et al., 2015). In October 2015, a severe haze episode was reported in the North China Plain (NCP), with the hourly peak concentration of particulate NO₃⁻ exceeding 70 μ g m⁻³ (Zhang et al., 2018b). Even in November 2018, during a heavy haze episode in northern China, the hourly peak concentration of PM_{2.5} still exceeded 289 μ g m⁻³, of which particulate NO₃⁻ accounted for 30% (http://www.mee.gov.cn/xxgk2018/xxgk/ xxgk15/201811/t20181116_674022.html, last access: 5 January 2019).

Another way to alleviate the particulate- NO_3^- pollution is to control NH₃ emissions. Previous studies were performed to demonstrate the necessity of NH₃ emissions abatement in reducing PM_{2.5} concentrations in the United States (Pinder et al., 2007, 2008; Tsimpidi et al., 2007; Wu et al., 2016) and Europe (de Meij et al., 2009; Bessagnet et al., 2014; Backes et al., 2016). Recently, a feature article pointed out that NH₃ could be key to limiting particulate pollution (Plautz, 2018). In contrast with low particulate matter pollution levels in the United States and Europe, what we are facing in northern China is the extremely high particulate- NO_3^- pollution, especially during severe winter haze events.

Although Fu et al. (2017) proposed that the NH₃ emission controls are urgently required in China, the effectiveness of NH₃ emissions mitigation to alleviate the particulate-NO₃⁻ peaks during severe winter haze episodes was seldom reported. Only Guo et al. (2018) used a thermodynamic model to estimate the sensitivity of particulate NO₃⁻ to TA (sum of ammonia and ammonium) during one winter haze episode in Beijing. In their study, the atmospheric chemistry simulations based on NH₃ emission controls scenario were lacking to demonstrate the regional effects.

It is urgent to alleviate severe particulate- NO_3^- pollution in northern China, and the study on the effectiveness on NH_3 emission controls is necessary. In this study, we firstly compile a comprehensive NH_3 emission inventory for northern China for the winter of 2015 and estimate the NH_3 emission reductions by improving manure management. Then, the thermodynamic equilibrium model (ISORROPIA-II) and Weather Research and Forecast Model coupled with chemistry (WRF-Chem) are used to investigate the effectiveness of NH_3 emission reductions in alleviating particulate $NO_3^$ during a severe haze episode. The molar ratio based on observations is used to explore the efficiency of particulate- $NO_3^$ reductions during the severe haze conditions in wintertime.

2 Methods and materials

2.1 Observational data

Hourly time-resolution aerosol and gas measurements were conducted at the Peking University urban atmosphere environment monitoring station (PKUERS) (39.991° N, 116.313° E) in Beijing in December 2015 and December 2016. A commercialized semicontinuous in situ gas and aerosol composition (IGAC) monitor was used to measure the concentrations of water-soluble ions (e.g., NH_4^+ , SO_4^{2-} , NO_3^- , Na^+ , K^+ , Ca^{2+} , Mg^{2+} , Cl^-) in $PM_{2.5}$ and inorganic gases (e.g., NH₃, HNO₃, HCl). Relative humidity (RH) and temperature were observed at 1 min resolution at the same site. The quality assurance and control for the IGAC was described in Liu et al. (2017b). A typical severe haze episode occurred from 6 to 10 December 2015, with daily average concentrations of $PM_{2.5}$ exceeding 150 µg m⁻³ for 3 days (PM_{2.5} data are from the China National Environmental Monitoring Centre). The average RH and temperature in this haze event were 60.9 ± 11.4 % and 276.5 ± 1.4 K. The south wind was dominant with wind speed mostly less than 3 m s^{-1} . The average concentrations of particulate NO₃⁻, NH₄⁺ and SO₄²⁻ were 39.8 \pm 14.7, 27.7 \pm 8.6 and $42.4 \pm 16.0 \,\mu g \, m^{-3}$, respectively. The ratios of particulate-NO₃⁻ concentrations to SNA (sulfate, nitrate and ammonium) were 36.5 ± 4.0 %.

2.2 NH₃ emission inventory

A comprehensive NH₃ emission inventory of northern China (including the six provinces mentioned above) in December 2015 at a monthly and $1 \text{ km} \times 1 \text{ km}$ resolution is developed based on our previous studies (Huang et al., 2012; Kang et al., 2016). This is a brief introduction to our inventory. More detailed descriptions and validation are found in our previous studies. Our NH3 emission inventory is a bottomup process-based and statistical model which considers a diverse range of sources, including both agricultural (livestock manure and chemical fertilizer) and nonagricultural sectors (traffic, biomass burning, etc.). According to our inventory, the estimated NH₃ emission amount in northern China was 100 kt in December 2015. The largest source was livestock waste (57 kt, 57.0 % of the total emissions), following by vehicle (12.2%), chemical industry (8.8%), biomass burning (5.4%), waste disposal (4.0%), synthetic fertilizer applications (2.4%) and other minor sources (9.1%). The proportion of chemical fertilizer is small due to the limited fertilization activity in winter. In the past few years, our inventory has been compared with many studies to prove its reliability. For example, the spatial pattern of NH₃ emissions calculated in our inventory agreed well with the distribution of the NH₃ column concentrations in eastern Asia retrieved from the satellite measurements of the Infrared Atmospheric Sounding Interferometer (IASI) (Van Damme et al., 2014). In particular, our estimation of livestock NH₃ emissions in China is comparable to the results of Streets et al. (2003) and Ohara et al. (2007).

Another method for estimating NH_3 emissions is the inverse modeling method, which provides top-down emission estimates through optimizing comparisons of model simulations with measurements. For example, Paulot et al. (2014) used the adjoint of a global chemical transport model (GEOS-Chem) and data of NH_4^+ wet deposition fluxes to optimize NH_3 emissions estimation in China. Zhang et

al. (2018a) applied the the Tropospheric Emission Spectrometer (TES) satellite observations of NH₃ column concentration and GEOS-Chem to provide top–down constraints on NH₃ emissions in China. Their estimates are 10.2 and 11.7 Tg a⁻¹, respectively, which are close to our results (9.8 Tg a⁻¹) (Paulot et al., 2014; Zhang et al., 2018a). The accuracy of this method relies on many factors, such as the accuracy of initial conditions, the emission inventories, meteorological inputs, reaction rate constants and deposition parameters in the chemical transport model. Errors in these parameters could cause biases in the top–down estimation of NH₃ emissions. In addition, measurements of NH₃ or NH₄⁺ used in this method, including surface and satellite date, are usually sparse in spatial coverage and have uncertainties, which will also affect the estimation of NH₃ emissions.

2.3 ISORROPIA-II and WRF-Chem models

The thermodynamic equilibrium model, ISORROPIA-II (Fountoukis and Nenes, 2007), being used to determine the phase state and composition of an NH₄⁺–SO₄^{2–}–NO₃⁻–K⁺– Ca²⁺–Mg²⁺–Na⁺–Cl⁻–H₂O aerosol system with its corresponding gas components in thermodynamic equilibrium, was used to investigate the response of particulate NO₃⁻ to NH₃ emission reductions. Using measurements of watersoluble ions, *T* and RH from PKUERS as inputs and only including the thermodynamic equilibrium, ISORROPIA-II can avoid the inherent uncertainty in estimates of emission inventories, pollutant transport and chemical transformation. In this study, ISORROPIA-II was run in the "forward mode" and assuming particles are "metastable" with no solid precipitates, which is due to the relatively high RH range observed during this haze event (RH = $60.9 \pm 11.4 \%$).

We assess the performance of ISORROPIA-II by comparing measured and predicted particulate NO_3^- , NH_4^+ and gaseous HNO₃, and NH₃. An error metric, the mean bias (MB), is used to quantify the bias (the description of MB is shown below Fig. S1 in the Supplement). The predicted particulate NO_3^- , NH_4^+ and NH_3 agree well with the measurements and the values of R^2 are 0.99, 0.94 and 0.84, respectively (Fig. S1). The MB is only 1.0, 0.3 and $-1.8 \,\mu g \, m^{-3}$, respectively. However, the model performs poorly on HNO₃, with an R^2 of only 0.06 and an MB of $-1.0 \,\mu g \, m^{-3}$. This is because particulate NO_3^- is predominantly in the particle phase (the mass ratio of particulate NO_3^- to the total nitric acid (TN = NO_3^- + HNO₃) was 99.2 ± 1.9 %), and small errors in predicting particulate NO₃⁻ are amplified in HNO₃ predicting. Since the MB of HNO₃ is much smaller than the observed particulate NO_3^- (39.8 ± 14.7 µg m⁻³) and NH_4^+ $(27.7 \pm 8.6 \,\mu g \,m^{-3})$, this bias has little influence on simulating the efficiency of particulate- NO_3^- reductions.

In the real atmosphere, changes in the level of TA $(TA = NH_4^+ + NH_3)$ can affect the lifetime of TN (Pandis and Seinfeld, 1990). This is because the gaseous HNO₃ has a faster deposition rate in the atmosphere than particulate

 NO_3^- , and reductions in NH_4^+ may prompt particulate- $NO_3^$ partitioning into the gas phase. In such a case, the concentration of TN would not remain constant but decrease. In order to consider these, we use WRF-Chem (version 3.6.1) to investigate the effect of NH₃ emission controls on particulate- NO_{3}^{-} formation on the regional scale. The simulations were performed for the severe haze event from 6 to 10 December 2015. The modeling domain covered the whole of northern China with horizontal resolution of 25 km and 24 vertical layers from the surface to 50 hPa. The initial meteorological fields and boundary conditions were taken from the 6h National Centers for Environmental Prediction (NCEP) global final analysis with a $1^{\circ} \times 1^{\circ}$ spatial resolution. The inorganic gas-aerosol equilibrium was predicted by the Multicomponent Equilibrium Solver for Aerosols (MESA) in WRF-Chem (Zaveri et al., 2005). The Carbon-Bond Mechanism version Z (CBMZ) photochemical mechanism and Model for Simulating Aerosol Interactions and Chemistry (MOSAIC) were used in this study (Fast et al., 2006). Anthropogenic emissions from power plants, industrial sites, residential locations and vehicles were taken from the Multiresolution Emission Inventory for China (MEIC; available at http://www.meicmodel.org/, last access: 5 January 2019).

The performance of WRF-Chem is evaluated by comparing measured and simulated NO₃⁻, NH₄⁺, SO₄²⁻ and TA. Specifically, the observed and simulated values are as follows: (1) NO₃⁻ – 39.8 ± 14.7 µg m⁻³ versus 39.1 ± 15.6 µg m⁻³; (2) NH₄⁺ – 27.7 ± 8.6 µg m⁻³ versus 26.5 ± 11.7 µg m⁻³; (3) SO₄²⁻ – 42.4 ± 16.0 µg m⁻³ versus 39.7 ± 20.8 µg m⁻³; and (4) TA – 34.6 ± 8.5 µg m⁻³ versus 32.1 ± 11.0 µg m⁻³. The MBs of these four species are –0.7, -1.2, -2.7 and -2.5 µg m⁻³, respectively. Simulated particulate NO₃⁻, NH₄⁺, SO₄²⁻ and TA approximately agreed with the measurements (Fig. S2). There are still some simulation biases that may affect the simulation of particulate-NO₃⁻ reduction efficiency. This is discussed in detail in Sect. 3.3.

3 Results

3.1 High-potential reduction in wintertime NH₃ emissions in northern China

Livestock husbandry accounts for the largest proportion of NH₃ emissions in northern China in winter (approximately 60%), which is mainly caused by poor manure management. There are three main animal rearing systems in China: free-range, grazing and intensive. On the one hand, the proportion of intensive livestock husbandry in China is only about 40%, far lower than that of developed countries (Harun and Ogneva-Himmelberger, 2013). As a result, the widespread free-range and grazing animal rearing systems contribute more than half of the total livestock NH₃ emissions due to a lack of manure collection and treatment (Kang et al., 2016). On the other hand, there were no relevant regulations about

the storage of manure for intensive farms in China in the past few decades. This causes most livestock farms to also lack necessary measures and facilities for manure collection and storage (Chadwick et al., 2015).

Due to the current poor manure management in China, the improved manure management may have great potential for NH₃ emission reductions from livestock husbandry (Wang et al., 2017). The improved manure management mainly includes three phases: in-house handling, storage and land application (Chadwick et al., 2011). For winter, the emission reduction measures mainly focus on in-house handling and storage, since land application mainly occurs in spring and summer. According to previous studies, for in-house handling, regularly washing the floor and using slatted floor or deep litter to replace solid floor could both reduce NH₃ emissions by more than 50 % (Groenestein and Van Faassen, 1996; Monteny and Erisman, 1998; Gilhespy et al., 2009; Hou et al., 2015). For storage, covering slurry and manure could reduce NH₃ emissions by about 50 %–70 % (Balsari et al., 2006; Petersen et al., 2013; Hou et al., 2015; Wang et al., 2017).

Based on the above research results, the livestock NH₃ emission reduction strategies applied in this study include the following steps. Firstly, the proportion of intensive livestock production was raised from 40 % to 80 % in our NH₃ emission inventory model. In our model, the animals in free-range and grazing animal rearing systems are assumed to live outdoors for half a day, and the improved manure management is only effective for indoor animals. Therefore, increasing the proportion of intensive livestock production is conducive to better manure management (Hristov et al., 2011). Secondly, the ratios of NH₃ emission reductions mentioned above were multiplied by NH₃ emission factors in two phases of manure management: a 50% reduction for in-house handling and 60 % (average value of 50 % and a 70 %) reduction for storage. With these measures, we estimate that the NH₃ emission factors for livestock in China could be comparable to those in Europe and the USA (shown in Table S1 in the Supplement). Meanwhile, our NH₃ emission model predicted that the livestock NH₃ emissions were reduced by 60 % (from 57 to 23 kt), causing an approximately 40 % reduction in total NH₃ emissions. Spatially, NH₃ emissions decreased significantly in Hebei, Henan and Shandong, where the livestock NH₃ emissions accounted for a large proportion of the total (shown in Fig. S3).

3.2 Simulations of NO₃⁻ reduction due to NH₃ emission controls

In the ISORROPIA-II simulation, a 40 % reduction in TA was used to reflect the effects of reducing NH₃ emissions by 40 %. This approach has been used in many previous studies (Blanchard and Hidy, 2003; Vayenas et al., 2005). However, in the real atmosphere, the reductions in NH₃ emission are not always equal to the reductions in TA due to the regional



Figure 1. A comparison of particulate nitrate (NO_3^-) between the base (blue line) and emission reduction cases (red line) simulated by the ISORROPIA-II model in this severe haze episode.

transmission. Their differences are discussed in the WRF-Chem simulation.

In this haze event (from 6 to 10 December 2015), the mean concentration of particulate NO_3^- decreased from 40.8 to $25.7 \,\mu g \, m^{-3}$ (a 37 % reduction). In addition, the peak hourly concentration of NO₃⁻ decreased from 81.9 to 30.7 μ g m⁻³ (a 63 % reduction) (shown in Fig. 1). The fundamental thermodynamic processes of TA reductions in decreasing particulate NO_3^- are explained below. Firstly, we found that NH_3 was readily available to react with HNO₃ in the thermodynamic equilibrium system because NH₃ was $6.6 \pm 3.8 \,\mu g \,m^{-3}$ while HNO₃ was only $0.4\pm1.1\,\mu g\,m^{-3}$. Secondly, almost all particulate NO_3^- condensed into the aerosol phase (the mass ratio of particulate NO₃⁻ to TN was 99.2 \pm 1.9 %) under such low temperature conditions $(276.5 \pm 1.4 \text{ K})$. Thirdly, the NH₃-HNO₃ partial pressure production (K_P) was as low as about 0.1 ppb² (calculated from ISORROPIA-II outputs, depending not only on temperature and RH but also on sulfate concentration). The value of K_P would remain constant if the temperature, RH and sulfate concentration remained unchanged. In general, NH₄NO₃ was not easy to volatilize into the gas phase under these circumstances.

When TA was reduced by 40 %, the average mass concentration of gaseous NH₃ decreased from 6.6 to $0.01 \,\mu g \,m^{-3}$ (from 8.8 to 0.05 ppb). In order to keep the value of K_P constant in the thermodynamic equilibrium state, the reductions in NH₃ increased HNO₃, which shifted the particulate-NO₃⁻ partitioning toward the gas phase. Hence, when NH₃ in gas phase was almost completely depleted, HNO₃ increased from 0.4 to 15.5 $\mu g \,m^{-3}$ (from 0.1 to 5.6 ppb), leading to a reduction in particulate NO₃⁻ from 40.8 to 25.7 $\mu g \,m^{-3}$ (a 37.0 % reduction). Meanwhile, NH₄⁺ also decreased from 27.9 to 20.6 $\mu g \,m^{-3}$, and there was almost no change in sulfate level (decreased from 39.7 to 39.3 $\mu g \,m^{-3}$), with only a trace amount of NH₄HSO₄ produced. This indicated that the



Figure 2. Spatial distribution of particulate- NO_3^- concentrations in northern China predicted by WRF-Chem from 6 to 10 December 2015, for (a) the base case, (b) the emission reduction case, and (c) the difference between the base case and the emission reduction case. The scope of this study focuses on the black box, including Beijing (BJ), Tianjin (TJ), Hebei (HB), Shanxi (SX), Shandong (SD) and Henan (HN).

reduction in particulate NH_4^+ and NO_3^- was mainly due to the reduction in NH_4NO_3 . The sum of particulate NO_3^- and NH_4^+ decreased from 68.7 to 46.3 µg m⁻³ (a 32.6 % reduction).

We also conducted WRF-Chem simulations to quantify the impacts of NH₃ emission controls on particulate NO₃⁻ regionally. A 60% reduction in livestock NH3 emissions was used as an emission reductions scheme, and Fig. 2 shows the spatial distribution of particulate NO_3^- under the base case and the emission reduction case. The spatial distribution of particulate NO₃⁻ was mainly concentrated in most parts of Henan (HN) and Hebei (HB), with average concentration over $30 \,\mu g \,m^{-3}$ (included in the black box shown in Fig. 2a). The highest particulate- NO_3^- concentrations, more than $60 \,\mu g \,m^{-3}$, were mainly located in central south of Hebei and northern Henan. In the emission reduction case, the mean concentration of particulate NO₃⁻ decreased from 30.6 to $18.5 \,\mu g \,m^{-3}$ (a 39.4 % reduction) in the range of the black box. Meanwhile, the particulate NH_4^+ decreased from 16.3 to 11.7 μ g m⁻³ (a 28.1 % reduction). The sum of particulate NO₃⁻ and NH₄⁺ decreased from 46.9 to $30.2 \,\mu g \,m^{-3}$ (a 35.6 % reduction). Besides, the sulfate concentration slightly changed from 19.7 to 17.6 μ g m⁻³, and PM_{2.5} concentration dropped from 143.4 to $125.4 \,\mu g \, m^{-3}$. The largest reductions in particulate NO_3^- were mainly located in the central north of Henan and central Hebei, where the percentage reduction was generally more than 60% (shown in Fig. 2b). In some areas with high particulate-NO₃⁻ concentrations, particulate NO_3^- had been effectively reduced by more than $30 \,\mu g \, m^{-3}$ (shown in Fig. 2c). In these regions, severe haze events occurred frequently due to their large emissions of air pollutants, including NH₃ (Wang et al., 2014; Zhao et al., 2017). In addition, TN was reduced by 34.1 % (from 31.8 to $21.0 \,\mu g \,m^{-3}$), which was in line with the assumption in Sect. 2.3. Correspondingly, TA decreased by 40.7 % (from 17.2 to $10.2 \,\mu g \, m^{-3}$), very close to the reductions in NH₃ emission (40%). This indicates that it is reasonable to use TA reductions to represent NH₃ emission reductions in the ISORROPIA-II simulation.

3.3 The particulate- NO_3^- reduction efficiency during the wintertime

The sensitivity of particulate NO_3^- to NH_3 is often determined by the availability of ambient NH_3 , which can be represented by the observable indicator (Seinfeld and Pandis, 2006). In this study, we use the observed molar ratio (*R*) of TA to the sum of sulfate, total chlorine and TN minus Na^+ , K^+ , Ca^{2+} and Mg^{2+} to represent the availability of ambient NH_3 and predict the sensitivity of the particulate NO_3^- to changes in TN and TA.

$$R = \frac{\text{TA}}{2\text{SO}_4^{2-} + \text{NO}_3^{-} + \text{HNO}_3(g) + \text{Cl}^{-} + \text{HCl}(g)} -2\text{Ca}^{2+} - \text{Na}^{+} - \text{K}^{+} - 2\text{Mg}^{2+}}$$
(1)

The accuracy of *R* was examined by constructing the isopleths of particulate- NO_3^- concentrations as a function of TN and TA (shown in Fig. 3). The NO_3^- concentration was constructed by varying the input concentrations of TA and TN from 0 to 200 µg m⁻³ in increments of 10 µg m⁻³ independently in ISORROPIA-II, while using the observed average value for the other components. Over a range of temperatures (273–283 K) and RHs (30 %–90 %), the dashed line of *R* = 1 divides each isopleth into two regions with a tiny bias, which indicates that *R* can be used to qualitatively predict the response of the particulate NO_3^- to changes in concentrations of TN and TA.

On the right side of the dashed line (R > 1), particulate-NO₃⁻ formation is HNO₃-limited. The NH₃ is surplus and almost all particulate NO₃⁻ exists in the aerosol phase. The TA reductions mainly reduce NH₃ with negligible effects on



Figure 3. Isopleths of the particulate-NO₃⁻ concentration (μ g m⁻³) as a function of TN and TA under average severe haze conditions in winter. The concentration of SO₄²⁻, Cl⁻, K⁺, Ca²⁺, Na⁺ and Mg²⁺ was 60.2, 9.3, 0.56, 0.04, 0.75 and 0.03 μ g m⁻³, respectively. Values are averages from all severe hazes during the observation period.



Figure 4. (a) The observed molar ratio (R) and the concentrations of SNA in PKUERS in December 2015 and December 2016. (b) The frequency of R during the same period.

particulate NO₃⁻. By contrast, particulate-NO₃⁻ formation is NH₃-limited to the left of the dashed line (R < 1). There is less NH₃ present in the gas phase, and TA reductions could reduce particulate NO₃⁻ efficiently. For example, when the concentrations of TN and TA are 100 and 50 µg m⁻³ (RH = 60 % and T = 273 K), the concentration of particulate NO₃⁻ is about 100 µg m⁻³ and the value of *R* is close to 1 (typical observational values during the severe haze in this study). In such cases, if TA were reduced by 50 % to 25 µg m⁻³, the particulate NO₃⁻ would be significantly reduced from 100 to 20 µg m⁻³, an 80 % reduction.

Under the typical winter conditions in northern China, the value of *R* was generally greater than 1 and gradually declining with the increase in SNA concentrations (shown in Fig. 4a). When the concentration of SNA is greater than $150 \,\mu g \, m^{-3}$, the values of *R* become close to and frequently lower than 1. This indicated that particulate-NO₃⁻ formation would easily become NH₃-limited under severe haze conditions when NH₃ emissions were reduced. In general, particulate NO₃⁻ will be reduced effectively by a 40% reduction in NH₃ emissions on the condition that the value of *R* is less than 1.4 (shown in Fig. S4). This situation accounts for 68.1% of all of December (shown in Fig. 4b). It should also

be noted that particulate NO_3^- is insensitive to a 40 % reduction in NH₃ emissions when the value of *R* is greater than 1.4 (shown in Fig. S4). This situation mainly occurs on relatively clean days (the concentration of SNA is less than 75 µg m⁻³), accounting for only 31.9 % of all of December (shown in Fig. 4a, b). Overall, reducing 40 % of NH₃ emissions could effectively reduce the levels of particulate NO_3^- under typical severe winter haze conditions in northern China.

The observed *R* provides a simple method to rapidly estimate the efficiency of NH_3 emission reductions in particulate- NO_3^- reductions, which can avoid the drawbacks of the air quality model, especially the uncertain estimates of meteorology. However, it also needs to be examined in more detail for specific pollution and meteorological conditions. Therefore, the observed indicator and air quality models should be used in a complementary way to assess the effectiveness of NH_3 emission control strategies.

Based on the above analysis, the influence of WRF-Chem simulation biases on particulate- NO_3^- reduction efficiency simulation mainly depends on the simulation bias of *R*. During the simulation case, the average simulated value of *R* is 1.3, which is equivalent to the observed value (1.3). Since WRF-Chem has a good estimation of the availability of ambient NH₃, its estimation of the efficiency of particulate- NO_3^- reductions is reliable.

It is noteworthy that the efficiency of particulate- NO_3^- reductions by NH₃ emission controls in northern China during severe winter hazes may be higher than that in the United States and Europe. Compared with our results (40% NH₃ emission reductions lead to about 40% particulate- NO_3^- reductions), in the United States and Europe, NH₃ emissions often need to be reduced by more than 70% before particulate NO₃⁻ begins to decrease (Pozzer et al., 2017; Guo et al., 2018). This is mainly because the strict emission controls of SO₂ and NO_x in these areas lead to a more ammonia-rich environment, which makes particulate NO₃⁻ insensitive to NH₃ emission reductions.

4 Conclusions

In this study, we found that during severe winter haze episodes, the particulate- NO_3^- formation is NH_3 -limited, resulting in its high sensitivity to NH_3 emission reductions. Meanwhile, livestock NH_3 emission control is a very efficient way to alleviate particulate- NO_3^- pollution during severe winter hazes. The estimations showed that improvements in the manure management of livestock husbandry could effectively reduce total NH_3 emissions by 40 % (from 100 to 60 kt) in northern China in winter. This would lead to a reduction in particulate NO_3^- by about 40 % (on average from 40.8 to 25.7 µg m⁻³) during severe haze conditions.

 NO_x emission controls could be a more direct and effective way to reduce particulate NO_3^- than NH_3 emission reductions. However, in northern China, the tar-

get of NO_x emission reductions is only about 25 % in the 13th Five-Year Plan (2016–2020) (http://www.gov.cn/ zhengce/content/2017-01/05/content_5156789.htm, last access: 5 January 2019). Due to the dominance of free-range animal rearing systems and the lack of emission controls policies, livestock NH₃ emission reductions in China could be practicable. In order to control PM_{2.5} pollution more effectively in northern China, measures to improve manure management in livestock urgently need to be implemented.

Data availability. The model input data and the NH₃ emission inventory used in this study are available from the corresponding author.

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Competing interests. The authors declare that they have no conflict of interest.

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