Elucidating the pollution characteristics of nitrate, sulfate and ammonium in $PM_{2.5}$ in Chengdu, southwest China, based on three-year measurements

Liuwei Kong[1], Miao Feng[2], Yafei Liu[1], Yingying Zhang[1], Chen Zhang[1], Chenlu Li[1], Yu Qu[3], Junling An[3], Xingang Liu[1,*], Qinwen Tan[2,*], Nianliang Cheng[4], Yijun Deng[5], Ruixiao Zhai[5], Zheng Wang[5]

[1]State Key Laboratory of Water Environment Simulation, School of Environment, Beijing Normal University, Beijing 100875, China

[2]Chengdu Academy of Environmental Sciences, Chengdu 610072, China

[3]State Key Laboratory of Atmospheric Boundary Layer Physics and Atmospheric Chemistry, Institute of Atmospheric Physics, Chinese Academy of Sciences, Beijing 100029, China

[4]Beijing Municipal Environmental Monitoring Center, Beijing 100048, China

[5]Yuncheng Municipal Ecological Environment Bureau, Yuncheng, 044000, China

* Corresponding author.

E-mail addresses: liuxingang@bnu.edu.cn (Xingang Liu) and 11923345@qq.com (Qinwen Tan)

Abstract

Nitrate, sulfate and ammonium (NSA) are the main secondary inorganic aerosols of $PM_{2.5}$ and play an important role in air pollution. In this study, a three-year observational experiment was conducted from January 1, 2015, to December 31, 2017, in Chengdu, southwest China. NSA pollution characteristics, chemical conversion generation, emission reduction control sensitivity and pollutant regional transport characteristics were analysed. NSA are the most important chemical compositions of particles with aerodynamic equivalent diameter ≤ 2.5 μm in ambient air ($PM_{2.5}$), and the contribution of nitrate to the accumulation of $PM_{2.5}$ concentration is greater than that of sulfate and ammonium. NSA also have obvious characteristics of annual, monthly, seasonal, diurnal and weekly variations. Through observation data and model simulation, it was also found that the existence of an aerosol aqueous environment plays

an important role in the formation and existence of NSA. Sensitivity analysis between NSAs found that controlling NO_3^- and SO_4^{2-} play an important role in reducing the contribution of NSA to $PM_{2.5}$, which also implies that the current control of NOx and SO_2 is important for improving air pollution. Combined with meteorological conditions and potential source contribution function (PSCF) analysis, local emissions and regional emissions of pollutants are found to have important impacts on Chengdu's atmospheric environment. This research result not only provides an assessment of the current atmospheric emission reduction effect but also provides an important reference for atmospheric pollution control.

Keywords: Secondary inorganic aerosols; Three-year measurements; Pollution characteristics; Chemical conversions; Regional transport; Chengdu

1 Introduction

In recent years, with the rapid development of China's domestic economy and acceleration of urbanization, energy consumption and pollutant emissions have also increased, which increases the burden on the atmospheric environment, and severe air pollution has become a focus of social concern (Liu et al., 2013a;An et al., 2019;Fu et al., 2014;Zhao et al., 2017). When air pollution forms, mass concentrations of particles with aerodynamic equivalent diameter ≤ 2.5 μm in ambient air ($PM_{2.5}$, also known as fine particles) can reach a higher pollution level, which not only reduces atmospheric visibility but also carries a large number of toxic species into the human lungs, increasing the risks of cardiovascular and cerebrovascular diseases (Chang et al., 2018;Tie et al., 2009;Kong et al., 2019;Zhao et al., 2018;Yang et al., 2015a). Nitrate, sulfate, ammonium, organic matter and elemental carbon are the main compositions of $PM_{2.5}$, among which nitrate, sulfate and ammonium (NSA) are the main secondary inorganic aerosols in $PM_{2.5}$ (Ji et al., 2019;Zheng et al., 2016). NSA mainly originates from the secondary aerosols produced by complex chemical reactions of NOx, SO_2 and NH_3 from coal combustion, vehicle exhaust emissions and agricultural sources (Liu et al., 2013b;Wang et al., 2016;Tian et al., 2017).

Because China's current main energy resource is still fossil fuels, which are widely used in industry, for vehicles and residentially, the emission reduction space of NSA is still restricted by a large number of gaseous precursors of NSA (Zhao et al., 2018;Tong et al., 2019). In addition, the chemical conversion of NO_2, SO_2 and NH_3 to form NSA is still very complex, and both homogeneous and heterogeneous reactions involve the chemical conversion of secondary inorganic aerosols, such as photochemical reactions, aqueous phase oxidation environments of aerosols and catalysis of mineral dust (Cheng et al., 2016;Sun et al., 2014;Wang et al., 2016;Ohta and Okita, 1990;He et al., 2014). The formation of sulfate can increase the acidity of aerosols (Sun et al., 2014). In contrast, the presence of NH_3 can play a role in neutralization and maintain the acid-base balance of aerosols (Wang et al., 2016). If improper control measures are taken in pollution reduction control, such as further ammonia emission reduction, the acidification of aerosols and environmental problems of acid rain may be aggravated (Liu et al., 2019c). In addition to the air pollution caused by the local emission of pollutants, the regional transportation of pollutants from its surrounding cities also has an important impact on the urban air quality. Determination of regional transport sources of pollutants, taking regional joint prevention and control measures, and jointly reducing the emissions of pollutants will enable better air control effects, particularly in the Beijing-Tianjin-Hebei region of northern China (Chen et al., 2019a).

Higher concentrations of NSA in $PM_{2.5}$ were also found in regions with more serious air pollution in China, such as Beijing-Tianjin-Hebei, the Yangtze River Delta, the Pearl River Delta, the Fenwei Plain, and the Chengdu-Chongqing region (An et al., 2019;Li et al., 2017;Liu et al., 2019d). In response to this situation, the Chinese government issued an Air Pollution Prevention and Control Action Plan (2013-2017) in 2013 to reduce pollutant emissions and improve air quality (the Sate Council, 2013, last access: June 17, 2020). A large number of treatment measures have been taken regarding coal combustion, motor vehicle emissions and outdated industrial capacities, and by 2017, China's ambient air quality control measures had achieved good results (Liu et al.,

2019a;Chen et al., 2019b;Cheng et al., 2019;Li et al., 2019a). In Beijing, $PM_{2.5}$, NO_2 and SO_2 decreased by 35.2%, 17.9% and 69.8%, respectively, in 2017 compared with 2013 (Beijing Municipal Ecology and Environment Bureau, 2018, last access: June 17, 2020). In Chengdu, $PM_{2.5}$, NO_2 and SO_2 decreased by 42.3%, 15.9% and 64.5%, respectively, in 2017 compared with 2013 (Chengdu Municipal Ecology and Environment Bureau, 2018, last access: June 17, 2020). To continue to promote air quality improvement, the Chinese government launched the "Three-Year Action Plan for Winning the Blue Sky Defense Battle" in 2018, which puts forward stricter requirements on how to further promote the implementation of emission reduction plans (the Sate Council, 2018, last access: June 17, 2020). Through long-term observations, a comprehensive analysis of $PM_{2.5}$ chemical compositions and source characteristics is carried out to verify the current implementation effects of emission reduction, and in-depth analyses of pollution reduction control characteristics are of great significance for the next step in air pollution control. However, these analyses may be affected by the experimental equipment, observation stations and other conditions, and the time span of these atmospheric observations usually includes several pollution processes or lasts for weeks or months. Thus, it is difficult to analyse the long-term variations in characteristics of air pollution through comprehensive observation. In particular, there are few high-time-resolution (1 hour) observation experiments carried out with online automatic observation systems (Sun et al., 2013;Tie et al., 2017;Guo et al., 2014). Especially in the Sichuan Basin of southwest China, there are few long-term observational experiments on NSA, which are the main chemical compositions of $PM_{2.5}$.

The Sichuan Basin is among the most important areas of air pollution in China (Qiao et al., 2019;Gui et al., 2019;Zhong et al., 2019). Although there are many studies in this area, there are few long-term studies of the hourly concentration data resolution of $PM_{2.5}$ chemical compositions. In this study, through three years of observations (from January 1, 2015, to December 31, 2017), we analysed the pollution characteristics of

NSA, as well as their formation mechanism and pollution control sensitivity. Finally, combined with local emissions and regional transport characteristics, we analysed the air pollution transport characteristics of Chengdu air pollution.

2 Experiment and methods

2.1 Observation site

Comprehensive observations were carried out at the Chengdu comprehensive observation station of atmospheric combined pollution (30.63°N, 104.08°E). The observation equipment was placed on the top of a building, approximately 25 m from the ground, and there was no obvious pollution source within approximately 200 m. The site is located in south section 1 of Yihuan Road, Wuhou District, Chengdu (Fig. 1), and traffic emission sources may be the main pollution emission source around the observation station. This is a typical residential, traffic and commercial mixed area that represents the characteristics of the urban atmospheric environment. Chengdu is also a megacity in the Sichuan Basin of southwest China, as well as an important part of the Chengdu-Chongqing region, which is among the regions with serious air pollution in China, and as shown in Fig. 1, the Sichuan Basin also has high aerosol optical depth (AOD).

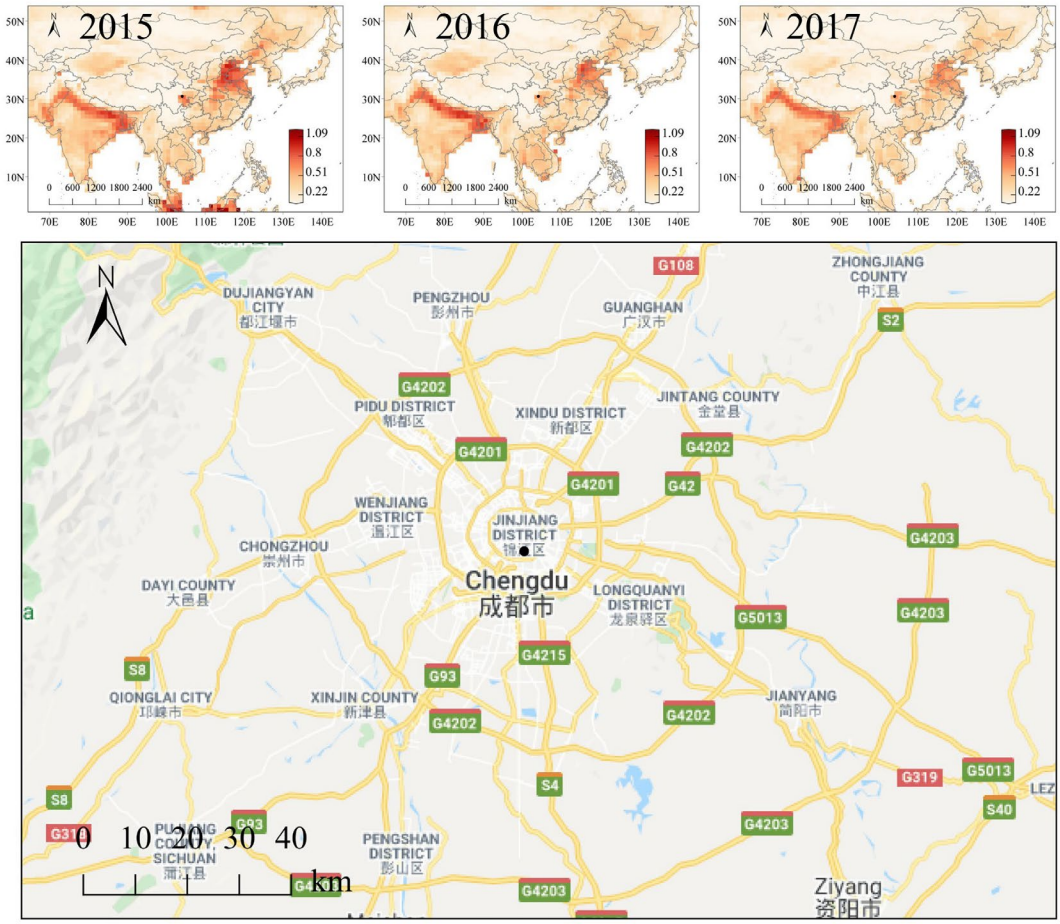


Fig. 1. Observation site in Chengdu. The image on the top shows the aerosol optical depth (AOD, 550 nm) from 2015 to 2017 (National Aeronautics and Space Administration, 2019, last access: June 17, 2020). The black dot in the image on the bottom shows the location of the observation site in Chengdu (background map from Google Maps, last access: June 17, 2020).

2.2 Instruments

During the research period, online experimental monitoring instruments were used to obtain the observation data with an hourly resolution (1 hour). The equipment list is shown in Table 1. Data quality control and assurance are an important part of the atmospheric comprehensive observation experiment, and this result is described in detail in the supplementary materials (Fig. S1-4).

Table 1. The experimental instruments used in this study

Instruments	Parameters	Manufacturer/Country
URG-9000	$NO_3^-/SO_4^{2-}/NH_4^+/Na^+/Mg^{2+}/Ca^{2+}/Cl^-/K^+$	Thermo Fisher Scientific/USA
SHARP 5030	$PM_{2.5}$	Thermo Fisher Scientific/USA
RT-4	OC/EC	Sunset Laboratory/USA
Xact-625	Metal elements	Cooper Environmental Services /USA
17i/450i/48i/49i	$NOx/NO_2/NO/NH_3/SO_2/CO/O_3$	Thermo Fisher Scientific/USA
WXT520	Meteorological parameters	VAISALA/Germany

OC: organic carbon; EC: element carbon

2.3 Chemical conversions and model methods

To clarify the conversion of gaseous pollutants to secondary aerosols, the nitrogen oxidation ratio (NOR) and sulfur oxidation ratio (SOR) were used to reflect the conversions of NO_2 and SO_2 to NO_3^- and SO_4^{2-}, respectively (Sun et al., 2014; Yang et al., 2015a). These ratios can be calculated using Eq. (1) and Eq. (2):

$$NOR=nNO_3^-/(nNO_3^-+nNO_2) \quad (1)$$

$$SOR=nSO_4^{2-}/(nSO_4^{2-}+nSO_2) \quad (2)$$

where n is the molar concentration.

The ISORROPIA-II thermodynamic model was used to analyse the variation in interaction characteristics among aerosol chemical compositions (Fountoukis and Nenes, 2007;Guo et al., 2017a;Ding et al., 2019). Temperature (T), relative humidity (RH) and the concentrations of Na^+, SO_4^{2-}, NH_3, NO_3^-, Cl^-, Ca^{2+}, K^+ and Mg^{2+} were input into the ISORROPIA-II thermodynamic equilibrium model. In this study, we used the "forward problems" mode to run the model, assuming that the aerosols were in a "metastable" state (salts do not precipitate under supersaturated conditions). At the time of data input, NH_3 data were the sum of NH_3 and NH_4^+. Previous studies had shown that the model has better performance when the RH is greater than 30%, and some studies also believe that the model performance is greater than 40%, so this study maintains the RH at higher than 40% when data are input (Ding et al., 2019;Guo et al., 2016). The simulated data and observed data were compared and analysed, and the

observation data of NH_3 were consistent with the input data of the model. The linear regression fitting slope of NH_3 was 0.96 (R^2=0.98), which showed that the run result of the model had good reliability and performance (Ding et al., 2019). Simultaneously, the aerosol water content (AWC) was calculated, and the sensitivity of the interaction between aerosol chemical compositions (NSA) and the pH of aerosols was analysed (Ding et al., 2019;Fountoukis et al., 2009). The pH was calculated using Eq. (3):

$$\mathrm{pH} = -\log_{10}\mathrm{H}^{+}_{\mathrm{aq}} \cong -\log_{10}\frac{1000\mathrm{H}^{+}_{\mathrm{air}}}{\mathrm{AWC}} \qquad (3)$$

where $\mathrm{H}^{+}_{\mathrm{aq}}$ (mol/L) is the concentration of hydronium ions in liquid water of atmospheric particulate matter, which can be calculated by the $\mathrm{H}^{+}_{\mathrm{air}}$ and AWC (μg/m^3) outputs from the ISORROPIA-II thermodynamic equilibrium model (Ding et al., 2019;Guo et al., 2017a).

2.4 CPF and PSCF methods

We used the conditional probability function (CPF) to analyse the characteristics of pollutants under the influence of wind direction (WD) and wind speed (WS). The analysis results using CPF were obtained using the R programming language, named openair. This function can be defined as CPF= $m_{\theta,j}/n_{\theta,j}$, where $m_{\theta,j}$ is the number of samples in the WD interval θ and WS interval j with mixing ratios greater than some 'high' pollution concentration, and $n_{\theta,j}$ is the total number of samples in the same WD-WS interval (Uria-Tellaetxe and Carslaw, 2014). Usually, a higher given 'high' pollution concentration (percentile) is chosen, such as the 90th percentile, which will mask the lower percentile pollution concentration source contributions. In this work, to obtain a more complete contribution of pollution sources, a range of percentile values, 0-25, 25-50, 50-75 and 75-100 were selected for the CPF calculation.

The potential source contribution function (PSCF) is based on an analysis of pollution sources given the airmass backward trajectory and can be used to judge the long-distance regional transport of pollutants (Ji et al., 2019). In this study, MeteoInfoMap and TrajStat (Wang et al., 2009) were used, and the model simulation data input model was provided by the National Oceanic and Atmospheric Administration (National

Oceanic and Atmospheric Administration, 2019, last access: June 17, 2020); these data were calculated to the 24-hour backward trajectories at the observation site at a height of 500 m every 1 hour from January 1, 2015, to December 31, 2017 (UTC+8). The calculated domain for PSCF was a range of 20-50° N, 75-115° E, and a grid cell with a resolution of 0.5°×0.5° was divided. The PSCF could be defined using Eq. (4):

$$PSCF_{ij} = \frac{M_{ij}}{N_{ij}} W_{ij} \quad (4)$$

$$W_{ij} = \begin{cases} 1.0 \left(N_{ij} \geq 3N_{ave}\right) \\ 0.7 \left(3N_{ave} > N_{ij} \geq 1.5N_{ave}\right) \\ 0.4 \left(1.5N_{ave} > N_{ij} \geq N_{ave}\right) \\ 0.2 \left(N_{ave} > N_{ij}\right) \end{cases} \quad (5)$$

where $PSCF_{ij}$ is the value for the ijth grid cell and M_{ij} is the total number of endpoints in the ijth grid cell, with pollution concentrations at the observation site (30.63°N, 104.08°E) that are greater than a given threshold value (the 75th percentile was selected for gaseous pollutants). N_{ij} is the number of backward trajectory endpoints in the ijth grid cell (0.5°×0.5°) during the simulation period. Therefore, the PSCF reflects the two-dimensional planar position distribution characteristics of potential sources, not the three-dimensional characteristics that reflect the transmission of pollution. To reduce the uncertainty in N_{ij}, the empirical weight function W_{ij} was introduced in Eq. (5), where N_{ave} is the average of N_{ij} during the simulation period (Ji et al., 2019;Zhang et al., 2017;Wang et al., 2009).

3 Results and discussion

3.1 Pollution characteristics of the interannual and entire observation periods

The annual average mass concentration of NSA and its proportion in $PM_{2.5}$ are shown in Table 2. The annual averages of $PM_{2.5}$ were 67.78, 71.88 and 59.68 μg/m^3, corresponding to 2015, 2016 and 2017, respectively. However, the pollution of $PM_{2.5}$ in Chengdu was much higher than the annual secondary guideline value (35 μg/m^3, Ambient air quality standards/GB3095-2012) and the World Health Organization annual guideline value (10 μg/m^3). The same $PM_{2.5}$ pollution problem was also a serious problem in Beijing and Nanjing (Ji et al., 2019; Zheng et al., 2019). The annual

average mass concentration of NSA also changed significantly, and the difference was large. The Mann-Whitney U test showed that the variation in NO_3^- was nonsignificant ($p > 0.05$), and SO_4^{2-} and NH_4^+ had obvious significance from 2015 to 2017 ($p < 0.05$), indicating that NO_3^- had not decreased significantly, and there was an increase in 2017 compared to 2015. SO_4^{2-} continued to decline, and NH_4^+ was also lower in 2017 than in 2016. Notably, SO_4^{2-} and NH_4^+ decreased significantly in 2017 compared with 2015, but the variation in NO_3^- was nonsignificant. Meanwhile, the annual averages of NO_3^-/SO_4^{2-} were 0.95, 1.02 and 1.45 for 2015, 2016 and 2017, respectively, indicating that the contribution of NOx emissions sources to $PM_{2.5}$ was increased compared with that of SO_2 emissions sources (Li et al., 2017; Wang et al., 2015). As shown in Table S1, from 2013 to 2017, the emissions of NO_2 in Chengdu were obviously higher than those of SO_2, but $PM_{2.5}$, NO_2 and SO_2 all decreased due to the implementation of the Air Pollution Prevention and Control Action Plan launched by the Chinese government and a more detailed pollution control plan launched by Sichuan Province. From 2015 to 2017, the measures taken by Sichuan Province in the coordinated reduction of multiple pollutants have been continuously strengthened, and the scope of management and control has been continuously expanded, for example, in the improvement of desulfurization, denitrification and dust removal technologies in key industries, from accelerated improvement in 2015 to deeper improvement in 2017. The process of eliminating small coal-fired boilers began in 2015 and was completed in 2017, when the ultra-low-emission coal-fired power plant transformation was promoted. In terms of vehicle emission control, we accelerated the elimination of "yellow label" vehicles (general term for gasoline vehicles with emission levels lower than the national I emission standard and diesel vehicles with emission levels lower than the national III emission standard when new vehicles are finalized) and “old vehicles” (the emission level does not meet the national stage IV emission standard) in 2015 and basically completed the elimination of "yellow label" vehicles in 2017. The quality supervision of oil products has also been improved, and non-road mobile machinery pollution

control requirements were proposed in the 2017 plan (The People's Government of Sichuan Province, 2015, 2016, 2017, last access: June 17, 2020). Compared with 2015, NOx and SO_2 decreased by 5.98% and 32.35%, respectively, in 2017, which shows that the treatment of NOx and SO_2 emissions has achieved remarkable results, of which the SO_2 emission reduction effect is the best, followed by that of NOx. The effect of this emission reduction is due to air pollution prevention measures, especially measures of "electricity instead of coal" and "natural gas instead of coal" (refers to increased use of electricity and natural gas in the residential sector to reduce coal combustion).

Table 2. Comparison of annual mass averages ($\mu g/m^3$) and proportions (%) for NSA (nitrate, sulfate and ammonium) from 2015 to 2017.

	$PM_{2.5}$	SO_4^{2-}	NH_4^+	NO_3^-	$NO_3^-/PM_{2.5}$	$SO_4^{2-}/PM_{2.5}$	$NH_4^+/PM_{2.5}$
2015	67.78	10.37	6.14	9.13	0.129	0.165	0.088
2016	71.88	8.53	6.16	9.27	0.123	0.133	0.089
2017	59.68	6.88	5.01	9.17	0.141	0.132	0.079

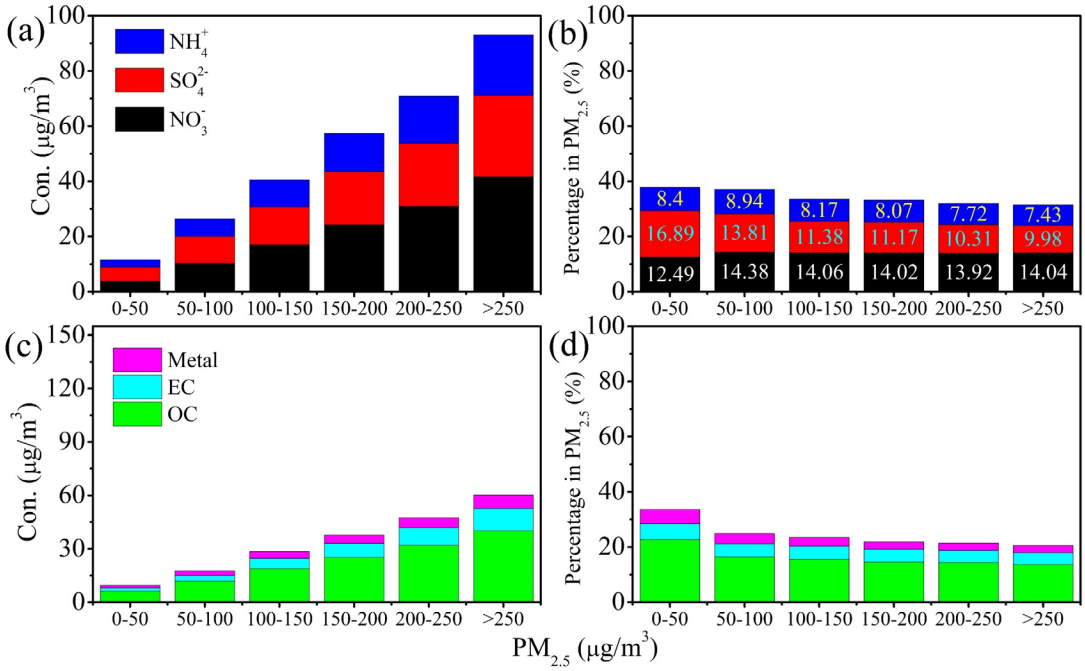


Fig. 2. Variation characteristics of the NSA (nitrate, sulfate and ammonium) and other chemical compositions with different concentrations of $PM_{2.5}$. (a) NSA mass

concentration. (b) Percentage of NSA in $PM_{2.5}$. (c) Chemical compositions of organic carbon (OC), elemental carbon (EC), and metal elements. (d) Percentage of OC, EC and metal elements in $PM_{2.5}$.

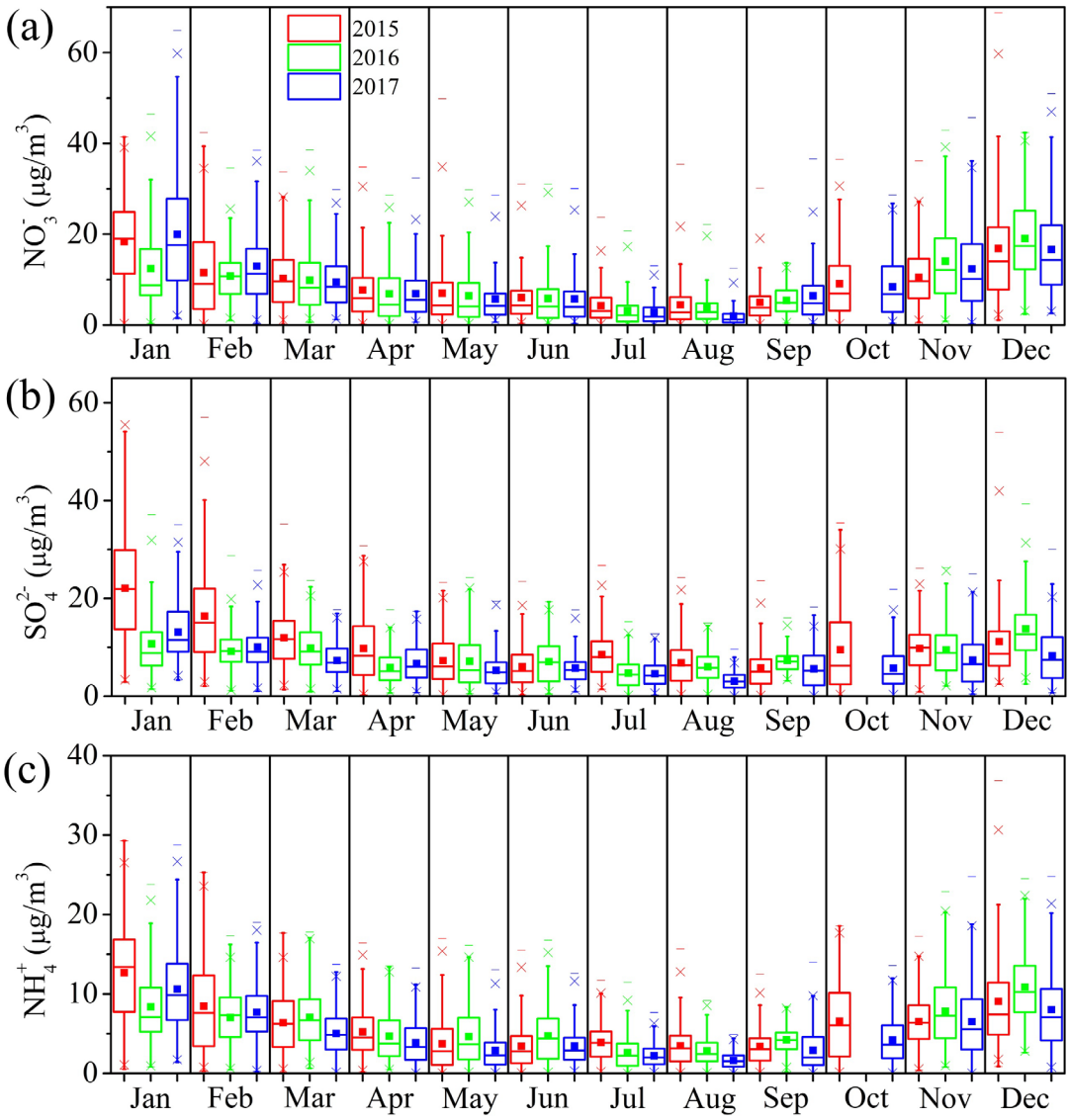
The chemical compositions of $PM_{2.5}$ from 2015 to 2017 varies with concentration, as shown in Fig. 2. With the accumulation of $PM_{2.5}$ in the atmosphere, the concentration of NSA also increased significantly, but the proportion of NSA in $PM_{2.5}$ decreased (Fig. 2a and b). The variation trend of OC, EC and metal elements with increasing $PM_{2.5}$ concentration is similar to that of NSA (Fig. 2c and d), and this variation trend of OC and EC is consistent with the results of long-term observation research carried out in Beijing (Ji et al., 2019). With the accumulation of $PM_{2.5}$ concentration, NSA, OC, EC and metal element concentrations have an increasing trend, but their ratio with $PM_{2.5}$ gradually decreases, indicating that other compositions have a higher contribution. This result, on the one hand, maybe since some chemical compositions such as ions and dust have not included in the statistics, on the other hand, the unknown component may also have a high contribution characteristic to $PM_{2.5}$ (Zhang et al., 2015;Huang et al., 2014). For the contribution characteristics of unknown components of $PM_{2.5}$, studies in some regions of China show that the contribution of higher $PM_{2.5}$ concentration is higher than that of lower $PM_{2.5}$ concentration (Huang et al., 2014;Li et al., 2017;Geng et al., 2019). When $PM_{2.5}$ was less than 50 $\mu g/m^3$ and greater than 250 $\mu g/m^3$, the mass concentrations of NSA were 11.57 and 90.06 $\mu g/m^3$, respectively, and the proportions were 37.78 and 31.45%, respectively. Comparing Fig. 2b and d, it was found that NSA was always the main contributor in the entire process of $PM_{2.5}$ accumulation, which was significantly higher than the proportions of OC and EC (Ji et al., 2019;Li et al., 2019b). In the accumulation process of $PM_{2.5}$ concentrations greater than 50 $\mu g/m^3$, NO_3^- accounts for a high proportion in NSA and is stable at approximately 14%, and the proportion of SO_4^{2-} and NH_4^+ continues to decrease (Li et al., 2019b;Wang et al., 2016). When the $PM_{2.5}$ concentration was less than 50 $\mu g/m^3$, the concentration of SO_4^{2-} was higher than that of NO_3^-, and the concentration of NH_4^+ was lower than the NH_4^+

concentration of $PM_{2.5}$ at 50 to 100 μg/m^3, possibly due to SO_4^{2-} concentration being higher than the NO_3^- concentration, forming more chemically stable $(NH_4)_2SO_4$ (Guo et al., 2017a). In addition, when the $PM_{2.5}$ was less than 50 μg/m^3, low RH and strong solar radiation were also important ways to generate sulfate (Yao et al., 2018).

3.2 Monthly and seasonal variations

The monthly variation characteristics of NSA from 2015 to 2017 are shown in Fig. 3. At the beginning and end of each year, the pollutant concentration is relatively high and relatively low in the middle of each year. The meteorological conditions also have obvious monthly variation characteristics (Fig. S5 a and b); from April to August, they have higher WS and lower RH, which is not only conducive to the dilution and diffusion of pollutants but also reduces the chemical conversions of pollutants by aqueous phase and influences the formation of secondary inorganic aerosols (Wang et al., 2016;Ji et al., 2019). Overall, the concentrations are higher in January and December and lower in July and August. The highest monthly average NO_3^- reached 19.98 μg/m^3 in January 2017, and the highest monthly average SO_4^{2-} and NH_4^+ were 22.08 μg/m^3 and 12.66 μg/m^3 in January 2015, respectively. The lowest concentrations of NSA appeared in August 2017, which were 1.96, 3.07 and 1.62 μg/m^3. The gaseous precursors of NSA also have obvious monthly variations, and the NOx and SO_2 trends were similar to those of NO_3^- and SO_4^{2-} (Fig. 3 and 4). NH_3 emissions were significantly different, with increases in warmer months (April-July) and colder months (September-December). On the one hand, NH_3 volatilization was promoted by relatively high Ts (Fig. S5c); on the other hand, the use of agricultural fertilizers and livestock farming were also important sources of NH_3 in China. Second, from urban region, fossil fuel combustion and motor vehicle emissions also contribute significantly (Liu et al., 2013b;Pan et al., 2016). Notably, NH_3 increased significantly from April to December 2017 compared with 2015 and 2016, especially during low-T months (Fig. 4c). The results of an analysis of the monthly concentration variation of pollutants indicate that the implementation of pollution reduction and control measures should be strengthened at

315 the beginning of each year (January to March) and the end of the year (October to
316 December).
317 The seasonal variation in NSA is shown in Fig. S6, and the concentration in winter was
318 much higher than that in summer. NO_3^- only declined in spring and summer from 2015
319 to 2017, with an increase in autumn and winter (Fig. S6a). Seasonal variations in NH_4^+
320 were similar to those of NO_3^-, with higher concentrations in winter and the lowest in
321 summer (Fig. S6c). This may be because higher Ts and WSs can not only promote the
322 decomposition of NH_4NO_3 in summer but also promote the dilution and diffusion of
323 pollutant concentrations (Guo et al., 2017a;An et al., 2019). There is a significant
324 downward trend in SO_4^{2-}, which continues to decrease in spring, summer and winter
325 from 2015 to 2017 (Fig. S6b). In autumn, the concentration was the highest in 2016,
326 and it was significantly lower in 2017 than in 2015 and 2016. The variation amplitude
327 of NSA and gaseous pollutants in cold months was significantly higher than that in
328 warm months (Figs. 3, 4 and S6). This higher variation amplitude may be due to the
329 differences in pollutant accumulation and scavenging processes. This finding also
330 indicates that the instability of local pollutant emissions and regional transport during
331 cold months was affected by meteorological conditions (Li et al., 2017;Ji et al., 2018).
332 The large variation amplitude of pollutants in different months, similar to the changes
333 in the Beijing-Tianjin-Hebei region of northern China and Chengdu, are due to the
334 accumulation and removal of pollution by meteorological conditions and pollutant
335 emissions (Ji et al., 2019;Qin et al., 2019;Zhang et al., 2019a).



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337 Fig. 3. Monthly variations in NSA (nitrate, sulfate and ammonium) concentrations from

338 2015 to 2017. (a) NO_3^-. (b) SO_4^{2-}. (c) NH_4^+.

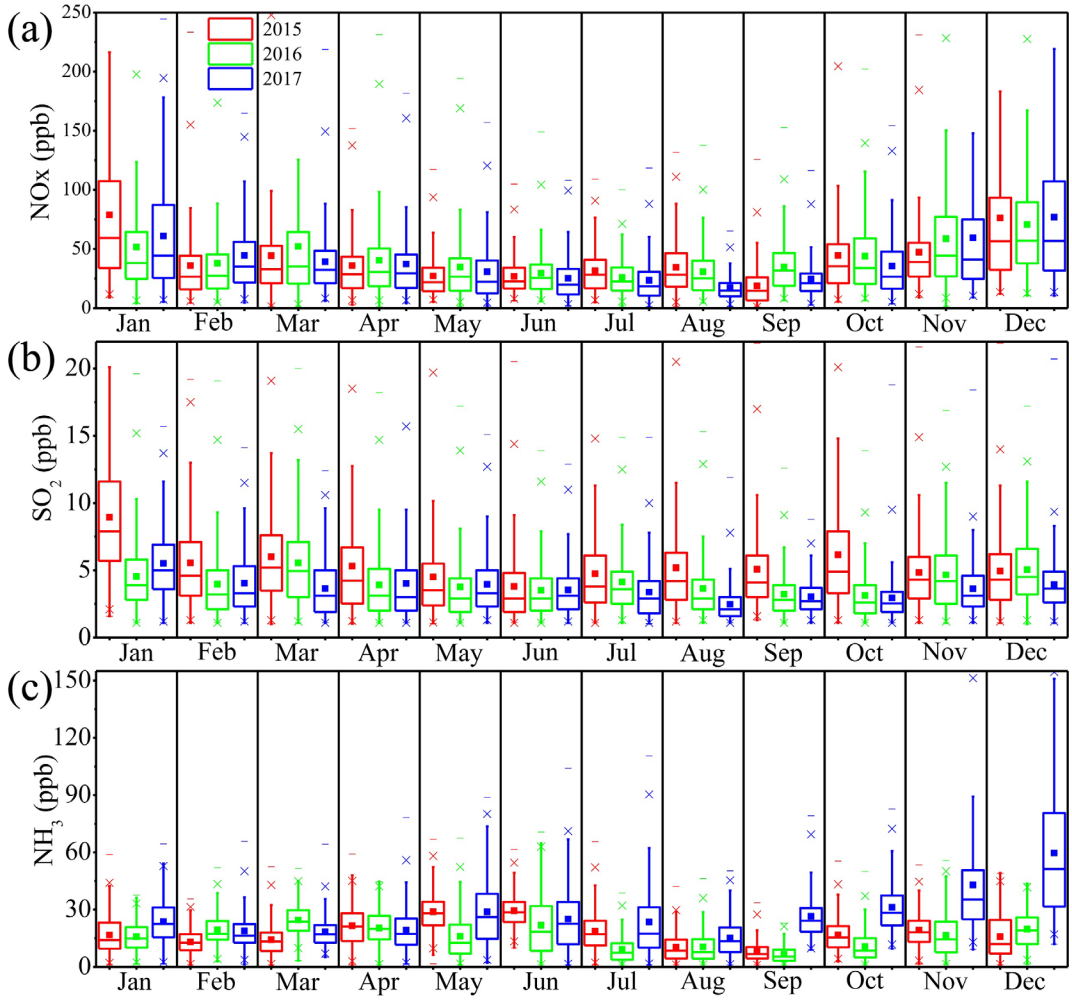


Fig. 4. Monthly variations in NOx, SO_2 and NH_3 concentrations from 2015 to 2017. (a) NOx. (b) SO_2. (c) NH_3.

3.3 Diurnal and weekly variations

From 2015 to 2017, the concentration of NSA was higher in the daytime than in the evening (Fig. 5a), and similar results were found in different seasons (Fig. 5b), which may be due to the combination of pollutant emissions and meteorological conditions. As shown in Fig. S7, from 9:00 to 11:00 a.m., the concentrations of SO_2, NOx, NH_3 and CO increased significantly, indicating that the primary emission of pollutants was relatively strong. At this time, although RH is in a declining stage, it still has a relatively high atmospheric humidity (approximately 65%), and O_3 and NO_2/NO also occasionally show an increasing trend, indicating that the atmospheric oxidizability has also increased (Figs. S7 and S8). This situation also provides favourable conditions for

the formation of secondary aerosols and promotes the accumulation of NSA (Cheng et al., 2016;Wang et al., 2016;Sun et al., 2014). In addition, before 10 o'clock, relatively low WS will enable easy pollutant concentration accumulation. In contrast, the higher WS in the afternoon may be the main factor for the decrease in pollutant concentration (Figs. 5 and S8). Photochemical reactions may also be one of the factors in the formation of NSA, and the concentration of O_3 peaks at approximately 15:00, which may be affected by the free radicals generated by photochemistry. At approximately 19:00, the ratio of NO_2/NO reached its highest value, and the concentration of NO_2 also increased significantly (Song et al., 2018;Zhu et al., 2019). At night, with the increase in RH (Fig. S8), dissolved ozone, free radicals, hydrogen peroxide and NO_2 can catalyse SO_2 to form secondary aerosols through an aqueous phase reaction (Zhang et al., 2015;An et al., 2019). The seasonal diurnal variation in NSA is shown in Fig. 5b. The concentration of NSA in winter was obviously higher than that in summer, and the diurnal variation range was larger. The concentrations in spring and autumn were closer, but the diurnal variation in spring was larger than that in autumn. The larger diurnal variation range not only indicates serious pollution but also indicates the importance of other factors affecting air quality, such as meteorological conditions and secondary aerosol conversion conditions (Ji et al., 2019;Yang et al., 2015b). The peak value of the NSA seasonal diurnal variation also varies in different seasons. The peak value appears at approximately 13:00 in winter, approximately 10:00 in spring and summer, and approximately 12:00 in autumn, possibly due to the influence of meteorological conditions. In previous studies in Beijing-Tianjin-Hebei and the Pearl River Delta, the concentration of pollutants was affected by meteorological factors, and it was usually lower in the daytime than at night. In the Yangtze River Delta, the peak usually occurs in the morning, but in our study, the concentration was higher in the daytime than at night (Peng et al., 2011;Wang et al., 2018;Guo et al., 2017b). In addition to the diurnal variations in WS and atmospheric humidity, some studies have shown that due to the unique topographical structure of the Sichuan Basin, the atmospheric circulation

between the Qinghai-Tibet Plateau, Yunnan-Guizhou Plateau and Sichuan Basin and the meteorological conditions of the Chengdu region are affected, such as the characteristics of air mass transport and typical “night rain” (more precipitation at night than in the day) under the influence of atmospheric circulation (Zhang et al., 2019c;Zhang et al., 2019b).

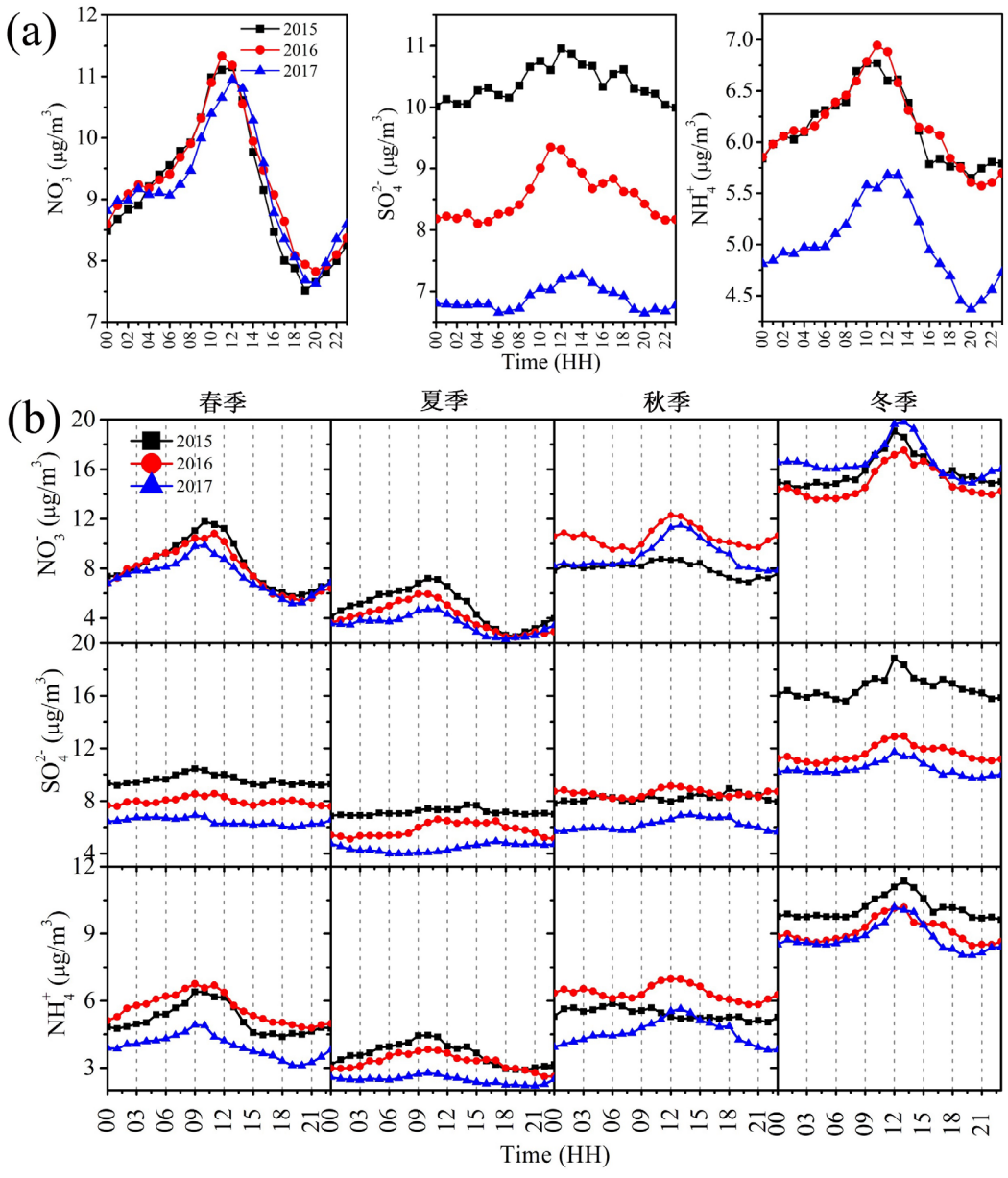


Fig. 5. Diurnal variations in NSA (nitrate, sulfate and ammonium) from 2015 to 2017. (a) Annual average. (b) Seasonal average.

The weekly variation in NSA is shown in Fig. S9. During the overall observational period, workdays (Monday to Friday) showed higher variations than weekends

(Saturday and Sunday), with the highest variation being on Tuesday and the lowest being on Sunday. Despite the difference in mean values between Tuesday and Sunday, nonparametric tests show that the difference in mean values was nonsignificant (Mann-Whitney U test, $P > 0.05$). As shown in Fig. S9, the average trends of NO_3^- and NH_4^+ were consistent from Monday to Sunday. The correlation coefficient was 0.94 ($P < 0.01$) from 2015 to 2017, which indicates that they have a common source and that vehicle emissions also have an important contribution to NH_4^+ (Pan et al., 2016). The average NO_3^-, SO_4^{2-} and NH_4^+ concentrations from 2015 to 2017 were 9.21, 8.64 and 5.64 ug/m^3 on workdays and 8.56, 8.33 and 5.29 ug/m^3 on weekends, respectively. The average values of NOx, SO_2 and NH_3 were 42.43, 4.35 and 20.39 ppb on weekdays and 39.60, 4.34 and 19.67 ppb on weekends, respectively. Similarly, the mean difference between NSA and gaseous precursors (NOx, SO_2 and NH_3) was not significant by the Mann-Whitney U test on weekdays and weekends. Population standard deviation comparisons of NO_3^-, SO_4^{2-} and NH_4^+ showed that workdays had higher standard deviations than did weekends, with 7.96, 6.04 and 4.35 on weekdays and 6.76, 5.69 and 3.88 on weekends, respectively, and it could also be seen from the box chart of NSA weekly variation that the concentration range on working days was slightly larger than that on weekends (Fig. S10). Analysis of the diurnal variation in NSA gaseous precursors on weekdays and weekends shows that the variation trend is relatively consistent (Fig. S11), and the concentration of NOx on weekdays will be slightly higher at the peak of 9:00 to 10:00 than on weekends, which may be affected by the morning rush hour of vehicles. In this study, NSA and gaseous precursors are also slightly higher on weekdays than on weekends, which indicates that in Chengdu's air pollution prevention and control actions, the management of relevant industries and departments should be strengthened on weekdays.

3.4 Chemical characteristics of NSA

3.4.1 Chemical conversion characteristics of NSA

Fig. 6 shows the abilities of NO_2 and SO_2 to chemically convert to NO_3^- and SO_4^{2-} at

different $PM_{2.5}$ concentrations. With the increase in $PM_{2.5}$ concentration, NOR and SOR gradually increased, indicating that the formation ability of NO_3^- and SO_4^{2-} increased during the formation of air pollution. In this study, when the $PM_{2.5}$ concentration was ≤ 50 μg/m^3, the average NOR and SOR were 0.07 and 0.27, respectively, and when the $PM_{2.5}$ concentration was greater than 250 μg/m^3, the average NOR and SOR increased to 0.22 and 0.41, respectively, indicating that the chemical conversion and formation ability of secondary inorganic aerosols was obviously enhanced when air pollution was aggravated. Previous studies suggested that when NOR and SOR were greater than 0.1 and 0.2, respectively, it has intense conversions and forms secondary inorganic aerosols (Yang et al., 2015a).

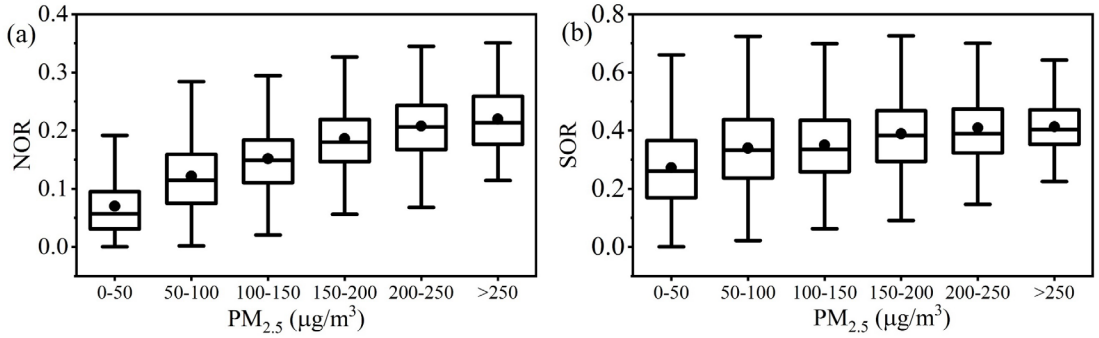


Fig. 6. Analysis of atmospheric chemical conversion ability at different $PM_{2.5}$ concentrations. (a) NOR (nitrogen oxidation ratio). (b) SOR (sulfur oxidation ratio).

Fig. 7 shows the variation characteristics of NSA chemical conversions with increasing RH. NOR and SOR increased with increasing RH, suggesting that NO_2 and SO_2 were more likely to produce NO_3^- and SO_4^{2-} under higher RH conditions. Previous studies have shown that the presence of NH_3 and NO_2 can promote the chemical conversion of SO_2 to SO_4^{2-} in the aqueous phase (Wang et al., 2016). In an aerosol aqueous phase environment, alkaline aerosol (NH_3) components can promote the dissolution of SO_2 and the formation of SO_4^{2-} under the oxidation of NO_2 (Cheng et al., 2016). Especially when the atmosphere is polluted, the formation of SO_4^{2-} by SO_2 through the aqueous phase environment can contribute most of the SO_4^{2-} (Sun et al., 2013). When the RH is greater than 80%, the NOR appears to decline, possibly because HNO_3 is semivolatile,

and the T increases at this time (Fig. 7 c), which is not conducive to the condensation of gaseous HNO_3 to particulate matter, which affects the amount of NO_3^- in $PM_{2.5}$ (Guo et al., 2017a). According to the ISORROPIA-II thermodynamic equilibrium model simulation, AWC also increases with RH (Fig. 7 d), and the increase in AWC can provide a liquid environment for aerosols, which is conducive to the dissolution and conversion of gaseous precursors of NO_2, SO_2 and NH_3 and promotes the formation of more NSA. The Pearson's correlation coefficients of RH and NOR and SOR were 0.12 and 0.16 ($p<0.01$), and the AWC and NOR and SOR were 0.73 and 0.37 ($p<0.01$), respectively, showing a significant positive correlation, indicating that the increase in AWC may be beneficial to the conversion of NO_2 and SO_2 to NO_3^- and SO_4^{2-}. As shown in Fig. S12, the simulated values of NSA (metastable state, liquid phase components) are compared with the observed data. The linear regression fitting slope is approximately 1 ($p<0.01$), indicating that the effect of the liquid phase environment in $PM_{2.5}$ is obvious; in addition, stable state simulation is also performed, and the linear regression fitting slopes of the NSA liquid phase state data output from the model and the observation data are 0.73, 0.63 and 0.74, and the Pearson's correlations are 0.82, 0.71 and 0.80 ($p<0.01$), indicating that they are more often combined with AWC in the aerosol aqueous phase environment at a stable state. Previous studies have also confirmed that the aqueous phase environment of aerosols plays an important role in the formation of secondary inorganic aerosols (Wang et al., 2016;Cheng et al., 2016).

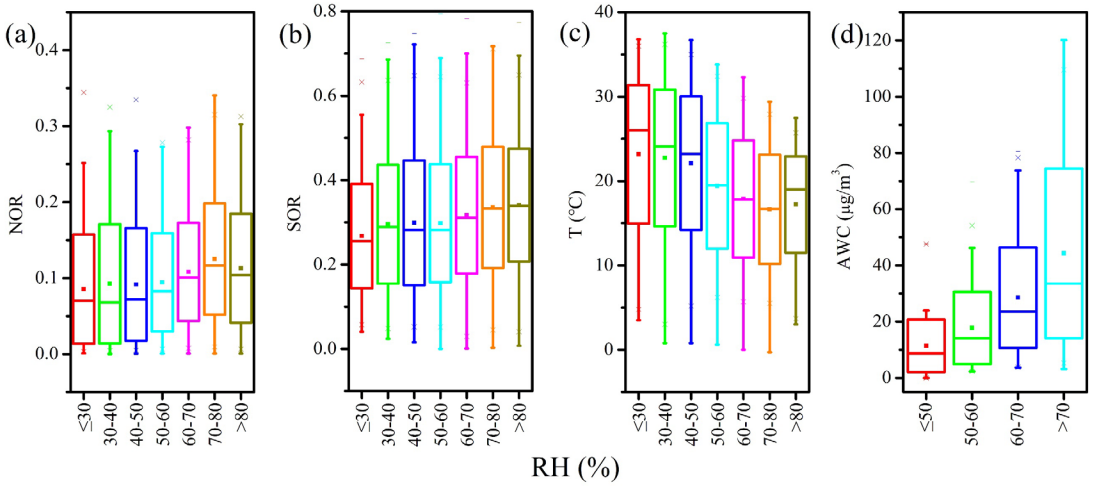
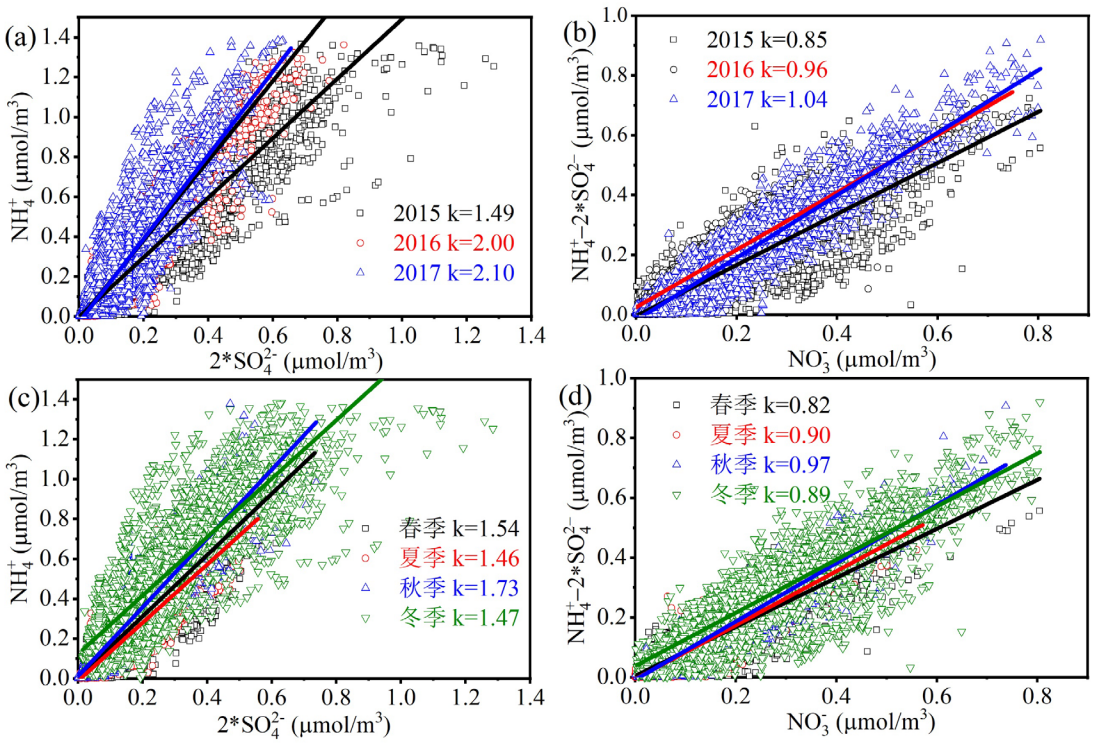


Fig. 7. Effects of RH on the chemical conversion of NSA (nitrate, sulfate and ammonium). (a) NOR (nitrogen oxidation ratio). (b) SOR (sulfur oxidation ratio). (c) Temperature (T). (d) AWC (aerosol water content).

3.4.2 Sensitivity analysis

The molar ratio analysis of NSA shown in Fig. 8 was used to analyse the chemical relationships among NSA. $(NH_4)_2SO_4$ and NH_4NO_3 are mainly composed of NH_4^+, SO_4^{2-} and NO_3^- in particulate matter (Malm and Hand, 2007;Meier et al., 2009). Because $(NH_4)_2SO_4$ has better stability than NH_4NO_3, NH_4^+ will first combine with SO_4^{2-} and then with NO_3^- (Liu et al., 2012). The annual average molar ratio of NH_4^+ to $2*SO_4^{2-}$ was more than 1, which indicates that SO_4^{2-} can be completely neutralized by NH_4^+ (Fig. 8a). The molar ratios of residual NH_4^+ (NH_4^+ - $2*SO_4^{2-}$) to NO_3^- were 0.85, 0.96 and 1.04 in 2015, 2016 and 2017, respectively. As shown in Fig. 8a and b, the gradual increase in the ratio (slope k) from 2015 to 2017 indicates that there is an increase in NH_4^+ in aerosol compared with SO_4^{2-} and NO_3^-, especially in 2017, with a ratio of 1.04, indicating the presence of other forms of NH_4^+, such as NH_4Cl and $(NH_4)_2C_2O_4$ (Sun et al., 2006). Seasonal variations in NH_4^+, SO_4^{2-} and NO_3^- are shown in Fig. 8c and d. The higher molar ratio in autumn indicates that the intensity of ammonia emission in autumn was higher than that in other seasons. This result also shows that the proportion of NH_4^+ relative to NO_3^- and SO_4^{2-} in $PM_{2.5}$ has increased. Therefore, while currently controlling NOx and SO_2 emissions, it is also necessary to strengthen NH_3 emissions control.



483

484 Fig. 8. Molar ratio analysis of NSA (nitrate, sulfate and ammonium). (a) Interannual
485 variation in the molar ratio of SO_4^{2-} and NH_4^+. (b) Interannual variation in the molar
486 ratio of NO_3^- and NH_4^+. (c) Seasonal variation in the molar ratio of SO_4^{2-} and NH_4^+. (d)
487 Seasonal variation in the molar ratio of NO_3^- and NH_4^+. k: Fitting slope of linear
488 regression.

489 Table 3 shows the sensitivity analysis of the concentration variations in SO_4^{2-}, NO_3^- and
490 NH_4^+. ISORROPIA-II thermodynamic equilibrium model sensitivity analysis is
491 described in detail in the Supplementary Materials. The coefficient of variance
492 represents the response of the species to variations in other components. The
493 coefficients of variance for NH_4^+ and NO_3^- produced by SO_4^{2-} changes were 52.22 and
494 1.70, respectively. Similarly, the coefficients of variance for NH_4^+ and SO_4^{2-} produced
495 by NO_3^- changes were 51.42 and 0.0005, respectively. The large coefficient of variance
496 for NH_4^+ indicates that the changes in NO_3^- and SO_4^{2-} can affect the presence of NH_4^+,
497 which also indicates that NH_4NO_3 and $(NH_4)_2SO_4$ were the main states of NH_4^+ (Liu et
498 al., 2012). The coefficients of variance for SO_4^{2-} and NO_3^- produced by TNH_3
499 (NH_3+NH_4^+) changes were 0.47 and 15.76, respectively, and the effect of TNH_3 on

SO_4^{2-} was less than that of NO_3^-, which indicates that NH_4^+ was excessive to SO_4^{2-} and that NH_4^+ first combines with SO_4^{2-} to form stable $(NH_4)_2SO_4$, and the remaining NH_4^+ and NO_3^- will combine to form NH_4NO_3.

Table 3. Sensitivity analysis of NSA (nitrate, sulfate and ammonium) concentration variations during the different observation periods.

Period	Variation	Coefficients of variance		
		NO_3^-	NH_4^+	SO_4^{2-}
2015-2017	NO_3^-		51.42	0.0005
	TNH_3	15.76		0.47
	SO_4^{2-}	1.70	52.22	

Coefficients of variance: standard deviation/average *100%;

Variation TNH_3: NH_3+NH_4^+ (μg/m^3);

Variation SO_4^{2-} and NO_3^- units: μg/m^3

Through the implementation of the Air Pollution Prevention and Control Action Plan, the reduction in SO_4^{2-} in $PM_{2.5}$ has achieved good results. Therefore, while continuing to promote "electricity instead of coal" and "natural gas instead of coal" to reduce coal combustion pollution, more stringent control measures should be added for NO_3^- and NH_4^+ emissions. To further improve air quality, the Chinese government launched a “Three-Year Action Plan for Winning the Blue Sky Defense Battle” in 2018 and proposed emission reduction targets for NOx and SO_2 emissions, which will be 15% lower in 2020 than in 2015 (the Sate Council, 2018, last access: June 17, 2020). The results of using the ISORROPIA-II thermodynamic equilibrium model to simulate NO_3^-, SO_4^{2-} and TNH_3 emission reduction control effects of 5%, 10%, 15% and 20%, respectively, are shown in Table S3, showing that controlling the concentration of NO_3^- and SO_4^{2-} is also helpful to reduce the concentration of NH_4^+ and indicating that controlling its precursor NOx and SO_2 is of great significance to reduce the secondary inorganic aerosol in $PM_{2.5}$ (the detailed results are described in the supplementary materials). Previous studies have also shown that the conversion of SO_2 to SO_4^{2-} in the

aqueous phase not only increases the conversion of SO_4^{2-} but also enhances the formation of NO_3^- in the aqueous phase (Wang et al., 2016). Therefore, SO_2 emission reduction may play a key role in the process of controlling emission reduction in NSA pollution, as it not only reduces the presence of NH_4^+ ($(NH_4)_2SO_4$) in particulate matter but also affects the formation of NH_4NO_3 by influencing the formation of NO_3^-. NO_2 and NH_3 can also promote the conversion of SO_2 to SO_4^{2-} through an aqueous phase environment (Wang et al., 2016). Therefore, priority control of NOx and SO_2 emissions is an important way to reduce NSA in particulate matter.

The increase in NSA can increase the hygroscopicity properties of aerosols, and more AWC can increase the pH by diluting the hydrogen ion concentration (Kong et al., 2020;Ding et al., 2019). Previous studies have also shown that SO_4^{2-} formation reduces aerosol pH (Sun et al., 2014). The effects of NO_3^-, SO_4^{2-} and TNH_3 on pH when using the ISORROPIA-II thermodynamic equilibrium model to simulate pollutant concentration reduction are shown in Table S3. With the decrease in NO_3^- and SO_4^{2-}, the pH value increases, but NO_3^- has no obvious effect on the pH value, SO_4^{2-} has an obvious effect on the pH value, which indicates that the formation of SO_4^{2-} in the aerosol can increase the acidity of the aerosol (Sun et al., 2014). The greater the reduction of TNH_3, the lower the pH value is, which shows that the presence of NH_3 as an alkaline gas can alleviate some of the acidity produced by SO_4^{2-} (Cheng et al., 2016). When the synergistic control of pollutants is reduced, it also has a certain effect on pH, increasing from 4.07 to 4.16. Some studies believe that if ammonia emissions are reduced significantly, the risk of acid rain may increase (Liu et al., 2019c). As shown in Fig. S13, the acid rain problem in China is mainly concentrated in southern China, especially in southwestern China, southern China, and the Yangtze River Delta in eastern China. Therefore, how to adjust the emission reduction ratio in combination with the characteristics of regional air pollution and energy consumption and thus help reduce the problem of aerosol acidity changes caused by air pollution reduction is a problem worthy of in-depth study. Therefore, when controlling NOx, SO_2 and NH_3

emissions, it is necessary to consider the aerosol acid and alkali changes caused by emission reduction.

3.5 Characteristics of local emissions and regional transport

3.5.1 Local emissions

The concentration of pollutants is obviously affected by meteorological conditions; for example, WS and WD can affect the accumulation and removal of pollutants (Li et al., 2016). Figs. S14-16 show the annual variation characteristics of NSA and gas precursors affected by the WS and WD using CPF. Overall, the higher WS was accompanied by a lower pollutant concentration. As the WS decreased, the pollution became serious, and the pollution hot spots were gradually concentrated. On the whole, when the WS was usually greater than 2 m/s, the pollution was light (pollutant concentration percentile was between 0-25). When WS was usually less than 1 m/s, the pollution was heavy (pollutant concentration percentile was between 75-100), which also reflects the distance and orientation between the emission source and the observation station, indicating that when the pollution was serious, the contribution of local source emissions was more prominent.

NO_3^- and NOx have similar distributions of pollution hot spots in the polar plot diagram (Fig. S14), and when the concentration percentile was 0-25, the hot spots were concentrated in the northeast and southeast directions and widely distributed. When the concentration percentile was 25-75, the sources of NO_3^- and NOx were distributed west，southwest and northeast of the observation site, and there were important contribution sources in the northwest direction (WS was approximately 3-4 m/s) in 2017. When the WS was approximately 1-2 m/s and the concentration percentile was 50-75, the important NOx source was in the northwest direction. When the accumulation of pollution concentration was high (concentration percentile was 75-100), the NO_3^- source was mainly concentrated in the east and southeast of the observation site, and NOx was distributed in the south and southeast, with WSs of less than 1 m/s. Additionally, the distribution of pollution hot spots was relatively wide in 2016 (the

annual mean values of NOx were 42.15, 43.99 and 39.63 ppb in 2015, 2016 and 2017, respectively), indicating that the source was relatively wide, which may be one of the reasons for the relatively high concentration. The SO_4^{2-} and SO_2 pollution sources affected by meteorological conditions also have similar distribution characteristics (Fig. S15). At a higher concentration of pollutants, the pollution hot spots of SO_4^{2-} were distributed in the east and southeast of the observation site, and SO_2 was distributed in the northeast, southeast and west. Compared with 2017 and 2016, the distribution of SO_2 pollution sources in 2016 was also more extensive, mainly in the west and northeast. The NH_3 emissions were slightly different from those of NOx and SO_2 (Fig. S16). Under conditions of high pollution concentration (concentration percentile was 75-100), the pollution hot spots were distributed in the west in 2015 (WS was approximately 2-3 m/s), in the north in 2016 (WS was approximately 1 and 3 m/s), and in the near distance in 2017 (WS was approximately 0.5 m/s). The higher pollution concentration was accompanied by a relatively higher WS (2015 and 2016), which indicates that the NH_3 emission transport in the surrounding area was more obvious, which may come from the surrounding agricultural source distribution area (Liu et al., 2019b;Liu et al., 2013b). The annual mean value of NH_3 emissions in 2017 was 27.91 ppb, which is significantly higher than those in 2015 and 2016 at 17.93 ppb and 16.55 ppb, respectively. During the 25-50 concentration percentile period of NH_3, there was a WS of approximately 2 m/s east of the observation site, and during the 50-75 concentration percentile period, there was an obvious source northwest of the observation site, with a WS of approximately 4 m/s. During the 75-100 concentration percentile periods, the pollution sources were mainly local. This shows that in 2017, in addition to the pollution sources being distributed in the east and northwest, the higher NH_3 emissions were also contributed by the surrounding emission sources northwest of Chengdu.

3.5.2 Gaseous precursors of NSA regional transport

The PSCF is used to analyse the potential source distribution of pollutants to determine the regional transport characteristics of pollutants (Ji et al., 2019). In addition,

604 considering the aerosol lifetime, SO_2 (approximately 9.6 d) and NOx (approximately 1
605 d) are also very different (Guo et al., 2014), and the research also shows that NH_3 is
606 significantly contributed by local source emissions (Walker et al., 2004). Therefore, we
607 comprehensively consider selecting a 24-hour backward trajectory to carry out PSCF
608 simulation in the Chengdu region. Fig. 9 shows the PSCF analysis of NOx, SO_2 and
609 NH_3, with significant differences in their potential source distributions. The higher
610 PSCF value of NOx was mainly distributed west, northwest and southwest of Chengdu
611 in 2015, northwest and south of Chengdu in 2016, and south, west and northeast of
612 Chengdu in 2017. The PSCF of NO_2 and NO (Fig. S17) also reflects that their potential
613 sources are mainly influenced by the interior of Sichuan Province, especially in the
614 cities around Chengdu. Chengdu is located along the western margin of the Sichuan
615 Basin, and it was also observed through satellite remote sensing data that the higher
616 NO_2 emissions in the Sichuan Basin were distributed in Chengdu and Chongqing (Fig.
617 S18a). The SO_2 emissions were widely distributed, mainly in the Sichuan Basin. Among
618 them, Leshan city and Meishan city south of Chengdu had higher SO_2 emissions, and
619 another higher emission source was distributed in Chongqing (Fig. S18b). The PSCF
620 analysis of SO_2 showed that the higher PSCF values were distributed in southern,
621 western and southwestern Chengdu. Therefore, a comparison of Figs. 9 and S18b shows
622 that the main source of SO_2 may be distributed in the southern, western and
623 southwestern margin region of the Sichuan Basin. In particular, Leshan, Ya'an and
624 Meishan were important potential sources. As shown in the PSCF analysis of NH_3 in
625 Fig. 9, the higher PSCF was also concentrated in the inner Sichuan Basin, especially in
626 the urban agglomeration around Chengdu. In 2015, the potential source of NH_3 was
627 mainly distributed in the southwest and northeast of Chengdu, with higher PSCF in
628 Nanchong and other regions. In the southwest, it was concentrated in Ya'an, Meishan
629 and Leshan. In 2016, potential sources were mainly distributed in the southwest of Ya'an,
630 Meishan, Leshan and the southern part of the Ganzi Tibetan Autonomous Prefecture.
631 There were two characteristics of potential sources in 2017. A relatively light source

was relatively close to Chengdu, and the high PSCF was in Chengdu, which can be considered the contribution of local emissions. The other contribution is obvious as a long-distance potential source contribution, mainly in some cities in the northeast, Nanchong, Guangyuan and Mianyang, and to a certain extent at the junction of Shaanxi, Gansu and Sichuan. In 2017, in addition to the contribution of local emissions, the contribution of regional transport in the northeast may also be an important reason for the higher NH_3 concentration. Fig. S19 shows satellite remote sensing data of NH_3. Overall, the higher NH_3 column concentration is distributed in the Sichuan Basin, mainly concentrated in the region near Chengdu, showing that NH_3 is more discharged in the Sichuan Basin, especially in the surrounding areas of Chengdu. In addition, through the analysis of the Multiresolution Emission Inventory for China (MEIC), it is also found that the higher NOx, SO_2 and NH_3 emissions in Sichuan Basin are mainly concentrated in Sichuan Basin, as shown in Fig. S20. It can also be seen that NOx is mainly concentrated in more-developed Chengdu and Chongqing, SO_2 emissions are obvious in Chengdu and Western Chongqing, NH_3 emissions are widely distributed, and there are higher emission characteristics in Chengdu and its surrounding areas. Therefore, according to the analysis of pollution emissions and PSCF in Chengdu, it is necessary to strengthen regional air pollution control and take regional joint prevention and control measures to reduce the impact of air pollutant regional transport.

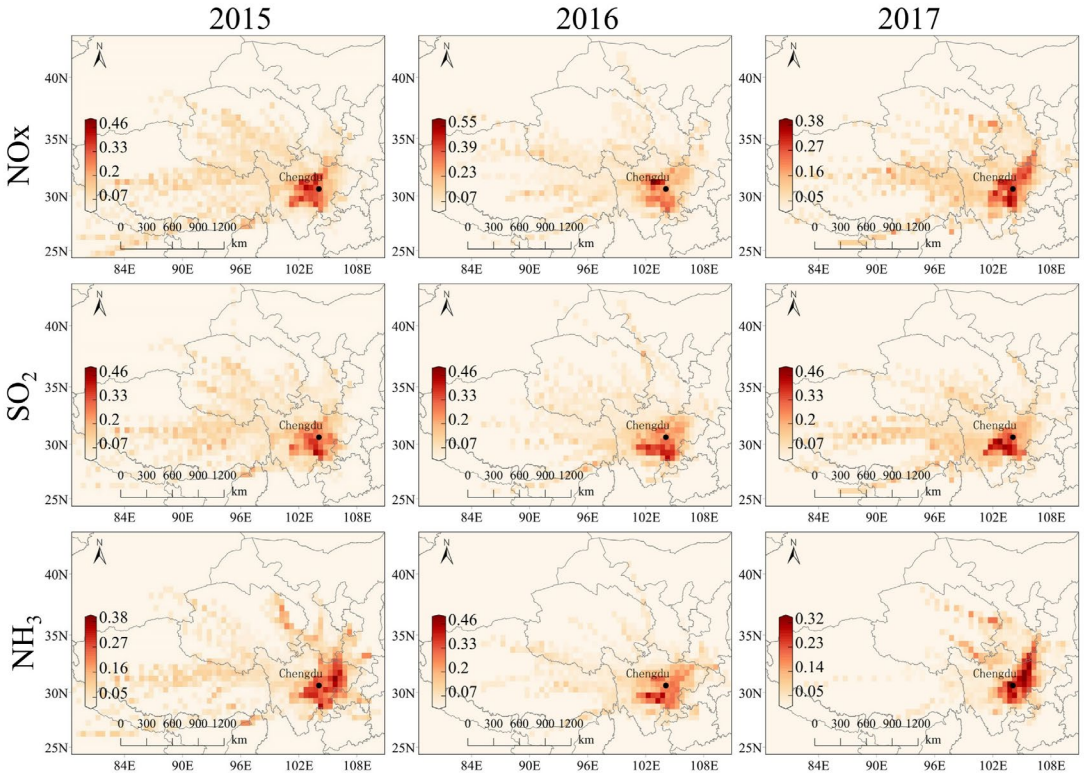


Fig. 9. PSCF (potential source contribution function) of NOx, SO_2 and NH_3 in Chengdu from 2015 to 2017.

4 Conclusions

The three-year observation experiment with hourly resolution of NSA from January 1, 2015 to December 31, 2017 was carried out in Chengdu in southwest China, which is in the Sichuan Basin. The pollution characteristics of NSA's annual, monthly, seasonal, diurnal and weekly variations were demonstrated. The characteristics of chemical conversion and the sensitivity of emission reduction control were analysed. Finally, combined with meteorological parameters and PSCF simulation, the local emission and regional transport characteristics of NSA gaseous precursors were also illustrated. The main conclusions were as follows:

(1) With the increase in $PM_{2.5}$ concentration, the NSA mass concentration increased, accounting for 31.45-37.78% of $PM_{2.5}$, and the contribution of NSA was higher than that of carbon aerosol (OM and EC). From 2015 to 2017, the contribution of NO_3^- to $PM_{2.5}$ increased, and in 2017, it became the main contribution component of NSA, and

it plays an important role in the concentration accumulation of $PM_{2.5}$. Higher and lower NSA concentrations were seen in winter and summer, respectively, and higher concentrations were seen more during the day than at night. Although the NSA concentration on weekdays was slightly higher than that on weekends, the mean difference between them was nonsignificant.

(2) With the increase in $PM_{2.5}$ concentration, there is an increasing trend of NOR and SOR, which indicates that the formation of NO_3^- and SO_4^{2-} increases obviously, and the increase in RH will promote the formation of NO_3^- and SO_4^{2-}. Using the ISORROPIA-II thermodynamic equilibrium model, it is found that NSA in aerosols is more likely to combine with AWC, which indicates that the aqueous environment of aerosols plays an important role in promoting the formation of NSA. The analysis of the interaction between NSA also confirmed that NH_4^+ will first combine with SO_4^{2-} to form $(NH_4)_2SO_4$, and the remaining NH_4^+ will combine with NO_3^- to form NH_4NO_3. The sensitivity analysis of NSA concentration shows that reducing NOx and SO_2 is beneficial to reducing NSA contribution in $PM_{2.5}$, but their changes also have an important impact on the pH of aerosols.

(3) Local emissions and regional transport of NSA gaseous precursors have an important impact on air pollution in Chengdu. When pollution is aggravated, the contributions of NOx and SO_2 to local emissions are relatively obvious. In addition to the local emission of NH_3, the contribution of pollution sources around Chengdu is also relatively obvious. PSCF analysis shows that the potential sources of pollution transmission in Chengdu are mainly distributed in Sichuan Province, and the most prominent contribution is made in Sichuan Basin, especially among the cities around Chengdu. The analysis of local emissions and regional transport shows that it is necessary to implement joint prevention and control of air pollution in the Sichuan Basin.

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Data availability

The data are available on request to the corresponding author.

Author contribution

XL, QT and LK designed and led this study. QT and MF were responsible for the observations. LK, MF, YL, YZ, CZ, and CL analysed the data. LK, YQ, JA, NC, YD, RZ and ZW discussed the results. LK and XL wrote the paper. All authors commented on the paper.

Competing interests

The authors declare that they have no conflicts of interest.

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