

The authors appreciate the reviewers for reviewing our manuscript and providing constructive comments. As suggested, we carefully revised the manuscript thoroughly according to the valuable advices, as well as the typographical, grammatical, and bibliographical errors. Listed below are our point-by-point responses in blue to the review's comments (in italic).

Anonymous Referee #2

This study analyzed concentrations of sulfate, ammonium, nitrate and other inorganic ions in precipitation and fine PM and estimated the fraction of the wet deposition attributed to in-cloud and below-cloud scavenging processes. The sequential sampling of precipitation over the course of a precipitation event provides interesting insight into changes in the precipitation chemistry that cannot be obtained from daily or weekly precipitation monitoring that are reported in many studies. The study introduced a newer approach to estimate the in-cloud and below-cloud wet scavenging proportions. In spite of the novel aspects in this study, the scientific discussions on wet scavenging of acidifying pollutants needs to be improved.

[Response]: We appreciate the valuable comments of the anonymous referee. We have prepared the point-by-point responses to address the reviewer's comments as shown below.

The results and discussion center on the wet scavenging of inorganic ions in PM_{2.5}. There was hardly any discussion on the wet scavenging of coarse particles and gas phase S and N compounds that can also contribute substantially to wet deposition of sulfate, nitrate and ammonium. The paper discussed relationships between precipitation and ambient air concentrations of inorganic ions; however, the latter was based on components in PM_{2.5}. How would the relationship change if coarse PM and gas phase compounds were included? Similarly for the below-cloud scavenging results, there needs to be a more balanced discussion on particle and gas-phase scavenging.

[Response]: Thanks for the comments. We are absolutely agreed with that the other sources, i.e., coarse particles and gas phase S and N compounds, can contribute to wet deposition of sulfate, nitrate and ammonium. Evenly, the model study in Japan by Kajino et al. (2015) showed the SO₂ and HNO₃ gases dominantly contributed to the washout of SO₄²⁻ and NO₃⁻, while the aerosol removal was dominated by the rainout process. However, in our measurement study, the on-line observation of coarse particles and gas phase S and N compounds were absent. We will try to address this in the revised manuscript and even in our future research.

1. The results in Kobe (high emission flux of SO₂ in Japan) showed the large difference between the observation and simulation, mostly due to the absent consideration of gas dissolved into raindrop in high-frequency observation measurements (Kajino et al., 2015). In their model set up, all of below-cloud gas SO₂ was assumed to be dissolved into raindrop and be fully oxidized to SO₄²⁻. However, as suggested by Seinfeld and Pandis (2006), the aqueous equilibrium

between ambient gas and precipitation cannot be assumed due to the relatively short residence times of falling precipitation. Considering the large amounts of particles ($60\text{--}90\text{ }\mu\text{g}/\text{m}^3$ in mass concentration) below-cloud in Beijing, the gases compounds may be not as important as that in simulation in Japan. According to the yearly book of “*Environmental Bulletin in Beijing*” from 1994 to 2017, the yearly concentration of SO_2 has a dramatically decreasing from $26.5\text{ }\mu\text{g}/\text{m}^3$ in 2013 to $8\text{ }\mu\text{g}/\text{m}^3$ in 2017. This relatively low-level concentration of SO_2 at surface may not contribute a dominant role in wet deposition of SO_4^{2-} . Similar case in NO_3^- , the ratio of gas-phase HNO_3 and the total NO_3^- in the summer in Beijing is only 0.12 according to the measurement study of Yue *et al.* (2013). We assumed the 10% ratio of gases added into the washout process, which only leads to less 5% difference of below-cloud scavenging contribution to total wet depositions. Anyway, for NH_3 , there might be larger uncertainties, since the high concentration of NH_3 at below-cloud layer over NCP (Pan *et al.*, 2017). Kasper-Giebl *et al.* (1999) reported that 49-79% of NH_4^+ in precipitation are from particulate ammonium, which indicate the large uncertainties of contribution from gases still exists in the form of NH_4^+ wet deposition. This also confirmed by our results: the large difference in the below-cloud contribution to NH_4^+ wet deposition estimated by the exponential approach and the average approach in Table 1 of the revised manuscript. Thus, the more research on the effect of NH_3 to NH_4^+ wet deposition in Beijing should be considered in the future.

2. As to particles wet scavenging, the main factors affecting the below-cloud scavenging include raindrop number size distribution, collection efficiency and raindrop terminal velocity. For collection efficiency, Brownian diffusion, directional interception, inertial impaction, thermophoresis and diffusion electrophoresis are the critical affected mechanisms. Coarse particles (aerosol particle sizes ranging from $2\text{--}20\text{ }\mu\text{m}$) are easily scavenged by inertial impaction. Especially coarse particles ($> 20\text{ }\mu\text{m}$) are also easily scavenged through the effect of gravity. Fine particles ($< 0.2\text{ }\mu\text{m}$) can be removed by Brownian diffusion. However, accumulation mode aerosols ($0.2\text{ }\mu\text{m}\text{--}2\text{ }\mu\text{m}$) are neither efficiently scavenged by Brownian diffusion nor by directional interception or inertial impaction. In addition, phoretic and electric charging effects mainly affected the particle size range of $0.2\text{--}3\text{ }\mu\text{m}$. And electrical effect is also one of undeniably mechanisms. Since the ions collected in precipitation are both from fine and coarse particles, the effect of coarse particles in wet deposition is considered in this study. However, as to the analysis of relationship between precipitation and ambient air concentrations of inorganic ions, the effects of coarse particles are not considered. This may bring little influence to SO_4^{2-} and NH_4^+ , but large uncertainties in NO_3^- , due to their particle size distribution characteristics. The lower coefficient in NO_3^- than SO_4^{2-} and NH_4^+ may also be attributed to this reason.

A more detailed explanation as to why the beginning and latter precipitation fractions represent below-cloud and in-cloud scavenging, respectively, is necessary since most of the results from this study are based on this key assumption. This data analysis

method seems to assume that below-cloud and in-cloud scavenging have to occur in sequence.

[Response]: Thanks for the comment. The data analysis method in this study do not assume that the below-cloud and in-cloud scavenging occur in sequence. Both of the below-cloud and in-cloud scavenging contribute to the total wet deposition (Seinfeld and Pandis, 2006). The general measurement to precipitation monitoring at ground surface cannot distinguish the two processes from each other as they are already mixed well in the raindrop falling down to the ground. However, according to (Seinfeld and Pandis, 2006), species can be incorporated into cloud and raindrops inside the raining cloud and this process determine the initial concentration of raindrops before they start falling below the cloud base. In this stage, despite of the efficient process of the nucleation scavenging in cloud, the total mass of aerosol in cloud is almost stable due to the slow process of interstitial aerosol collection by cloud droplets which is the determination process to aerosol mass. That is to say, the initial concentration of raindrops in cloud is well mixed and can be considered as a stable statue during the whole rainfall event. Actually, many observations in different regions (Aikawa *et al.*, 2009; 2014; Wang *et al.*, 2009; Quyang *et al.*, 2015; Xu *et al.*, 2017) reported that the chemical components in a rainfall event show a decayed variation with the increase of precipitation amount and eventually tends to a stable and low concentration level. The assumption in this study as well as the previous studies “*the concentrations in later increments can be attributed to scavenging by rainout only*” is based on this fact. It does not mean the below-cloud and in-cloud scavenging occur in sequence. But, instead, the two processes have been mixed in all stage of the rainfall event with the below-cloud scavenging contributed more in beginning fraction and the in-cloud scavenging contributed more in the later fraction due to the depletion of the air pollutants below cloud by washout. The question is, how we recognized the “later fraction” in a rainfall event. The 5 mm accumulated precipitation (the concentration of chemical components at the 5th fraction) was used in the previous studies. However, in many cases, the concentration of chemical ions after 5 mm accumulated precipitation cannot be reached at stable level due to different precipitation intensity and the concentration level of air pollutants in each rainfall event. In this study, the asymptote value from the exponential decay fitting curve of the observed rainwater concentrations was employed as the in-cloud ion concentration. To investigate the uncertainties of this approach, the comparison between the exponential approach and the average value in fractions 6 to 8 (here after, average approach) has been carried out. The results show the exponential approach gives lower estimates of in-cloud concentrations than the average approach, with the latter being recognized as an upper limit for in-cloud concentrations. The more detailed explanation on the estimation of below cloud scavenging has been added in the revised manuscript.

The discussion on temporal variations in precipitation concentration of inorganic ions is lacking some depth. Further analysis with annual emissions data are necessary to explain interannual changes and appreciate the impact of the air pollution control policies implemented.

[Response]: Agree. Further analysis on the temporal variations in precipitation

concentration of inorganic ions with change of annual emissions have been discussed in the revised manuscript. More detailed description can be seen in the response to the comments “L185” and “section 3.1”.

The way the below-cloud percentages are reported in the paper raises some critical questions. In this version, only the median below-cloud percentage were reported. It is uncertain how representative are the median values considering that each precipitation event is unique in terms of rainfall intensity, precipitation form (rain, snow, or sleet), air concentrations of chemical components, type of weather system, etc. It is important to state or discuss under what conditions apply to those below-cloud percentages and their limitations. A range of the below-cloud percentages should also be included as well as some discussion on the variability of the below-cloud scavenging percentages.

[Response]: Thanks for the comment. Indeed, each precipitation event is unique in terms of rainfall intensity, rainfall type, air concentrations of chemical components, etc. The average or median values for in-cloud concentration during 4 years cannot represent each unique precipitation event. In this study, the fitting curve is implemented in each rainfall event during 2014-2017. The median below-cloud percentage in Figure 1 is used to compare with the method that reported in previous studies (Aikawa et al., 2009;2014; Xu et al., 2017), in which the median value after 5 mm accumulated precipitation over year-round measurements were recognized as the in-cloud concentration. The comparison between the exponential approach and the average approach (average value after 5th fraction, i.e., 6th-8th) has been carried out. The results show the exponential approach gives similar ratio as the average approach for most ions, except for NH₄⁺, F⁻, K⁺ and Mg²⁺, where the maximum difference is less than 20% (Table 2). In general, the exponential approach gives lower estimates of in-cloud concentrations than the average approach. The latter may be recognized as an upper limit for in-cloud concentrations.

Based on this uncertainty analysis, the exponential approach to each unique event was made. Yearly ratio of below-cloud contribution to total wet deposition were calculated according to Eq (1-2), in which the \bar{C} is based on the median value of in-cloud concentration in all events in each year. This is consistent with that reported in (Aikawa et al., 2009), who also used the median value as the yearly in-cloud concentration. Besides, the estimations based on the average value as well as the first quartile and the third quartile are also included in the Figure 4 of revised manuscript. It is shown that the overall trend is the same as the median value. Thus, the conclusion is more robust through comparing with the different \bar{C} values.

To avoid the misunderstanding, the clarify of the method for estimating the below-cloud proportion has been added in the revised manuscript, which is as: *Noted that, the fitted exponential curve is applied to the combination of all 69 full events to estimate the yearly median concentration of chemical ions in-cloud and to compare with the results from previously reported method (i.e., median concentration after 5 mm increments). Besides, the exponential approach to each unique event was also employed.*

$$\text{Wetdep}_{\text{below-cloud}} = \sum_{i=1}^n (C_i - \bar{C}) \times P_i \quad (1)$$

$$\text{Contribution}_{\text{below-cloud}} = \frac{\text{Wetdep}_{\text{below-cloud}}}{\sum_{i=1}^n C_i \times P_i} \quad (2)$$

Besides, the sentence in L302 of the original manuscript has been revised as: *The ratio of below-cloud contribution to the four major components based on the yearly median value of the in-cloud concentration is also shown in Figure 4 in the revised version.*

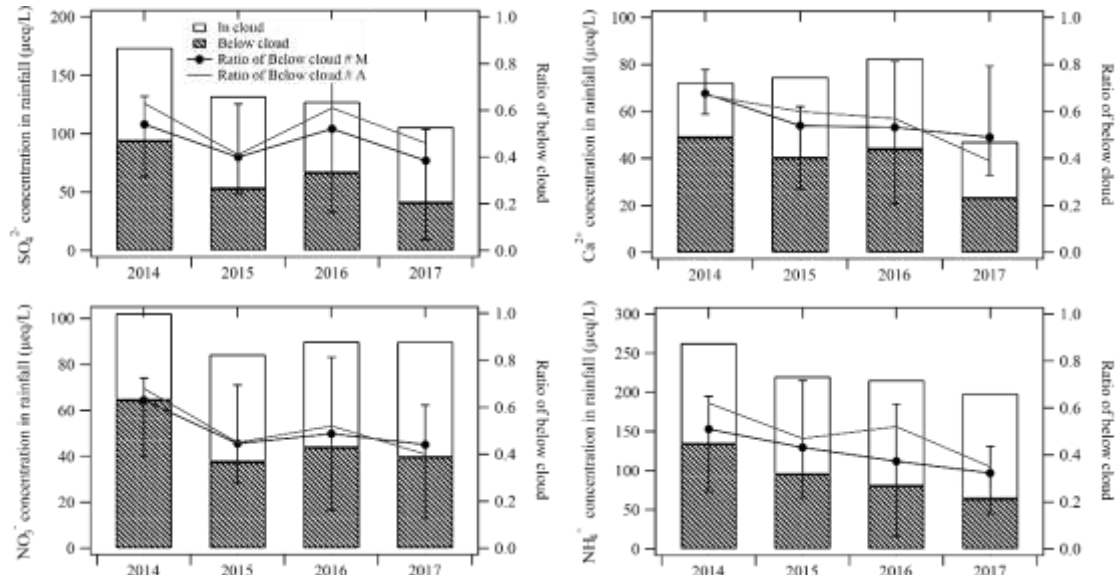


Figure 4 (in the revised manuscript). The annual volume weighted average below-cloud and in-cloud portion of SO_4^{2-} (a), Ca^{2+} (b), NO_3^- (c), and NH_4^+ (d) during 2014-2017. The ratio of annual median below-cloud contribution for each component is represented as the black line in each panel. The mark #M and #A in the ratio of below-cloud represent the estimation based on the median value and average value of in-cloud concentration in each year, while the first quartile and the third quartiles are also included in the figure.

Specific comments:

L22-24: Why would the Action Plan result in declines in Ca^{2+} given that it is mainly derived from crustal emissions?

[Response]: The Action Plan launched in 2013 including emission reduction not only from energy consumption of industry but also the raised dust in cities. This should result the decline in Ca^{2+} . Unfortunately, both of the emission and the concentration data of Ca^{2+} are absent in this study. Instead, the different of PM_{10} and $\text{PM}_{2.5}$ ($\text{PM}_{10} - \text{PM}_{2.5}$) air concentration during 2013-2017 have been calculated to represent the coarse particles, which contains the Ca^{2+} compound. The result shows that the concentration decreased from $31.2 \mu\text{g}/\text{m}^3$ in 2013-2014 to $24.0 \mu\text{g}/\text{m}^3$ over 2015-2017. This indicates the improvement of coarse particles even which is derived from crustal emissions have been made through the Action Plan launched in 2013 in China.

L25: “An improved sequential sampling method: : :” This is vague. The improvement to the method needs to be clarified in the abstract. It is not the improvement to the

sequential sampling method that is described in the paper, but rather the improvement to the method of estimating the below-cloud scavenging proportion.

[Response]: Thanks for the comment. The sentence has been revised as: *An improved method of estimating the below-cloud scavenging proportion based on sequential sampling is developed and implemented to estimate the contribution of below-cloud and in-cloud wet deposition over the four-year period.*

L27: Suggested revision, “below-cloud scavenging accounts for one half to two thirds of wet deposition”. To be more concise, only the second part of the sentence is necessary because it is very similar to the first part of the sentence.

[Response]: Agree. Considering the uncertainties of below-cloud scavenging contribution to NH_4^+ wet deposition, the sentence has been revised as: *Overall, the below-cloud scavenging plays a dominant role to the wet deposition of four major ions at the beginning of the Action Plan.*

L30: Was there a continuous year-to-year decline in the percentages, i.e. in 2015 and 2016? If so, this should be stated. If not, the next sentence, which suggests the result is due to the Action Plan, should not be included in the abstract because there is no clear decreasing trend.

[Response]: Thanks for the comment. No, there is not a continuous year-to-year decline in the percentages. The percentages from 2015 to 2016 show a little increasing change. Nevertheless, a decreasing trend, on the whole, have been observed in S and N compound both in airborne and precipitation after the Action Plan launched in 2013. All of the ions in precipitation except NO_3^- show a significant decrease ($p > 0.01$). Thus, the expressions on the Action Plan in the next sentence are kept in the abstract.

L36-37: “: : clearly identifies oxidized nitrogen species as a major target for future air pollution controls.” This statement seems to be based only on the lack of declining trends in airborne and precipitation nitrate. The observed concentrations of nitrate also need to be considered compared to other N or S compounds. In precipitation, it seems that the VWA of nitrate was similar to that of sulfate and ammonium was even higher than nitrate. It is not clear why the policy recommendation is to target oxidized nitrogen and not other N or S compounds as well.

[Response]: Thanks for the comment. We agree with that the other N and S compounds should be paid continuous attention in the future. However, the decreasing from both VCD NO_2 and nitrate is absent, while declining for S is significant. Besides, the statement is not only based on the lack of declining trend in airborne and precipitation nitrate, but also the decreasing ratio of S/N in both precipitation ($\text{SO}_4^{2-}/\text{NO}_3^-$, $\mu\text{eq/L}$) and air ($\text{SO}_4^{2-}/\text{NO}_3^-$, $\mu\text{g/m}^3$) as well as the decreasing ratio of $\text{NH}_4^+/\text{NO}_3^-$ from 2014 to 2017. These evidences clearly confirm that nitrate should be the major target for air pollution controls in the next Action Plan.

L44-46: Do these factors affect below-cloud rain scavenging only or both rain and snow?

[Response]: Yes, these factors affect both of the rain and snow through below-cloud scavenging. However, in this study, the measurement is only focused on rain instead of snow. The sentence has been revised as: *The below-cloud scavenging process depends both on the characteristics of the rain (snow).*

L57-58: For which chemical components do these results apply to? Can you provide details on the modeled and measured scavenging coefficients?

[Response]: Yes, we can provide details on the modeled and measured scavenging coefficients. Besides, the below-cloud scavenging coefficients used in CTM is referred to PM_{2.5}, while in observations is referred to SNA. The expression here is revised as: *They found that below-cloud scavenging coefficients for PM_{2.5} widely used in CTMs ($\sim 10^{-5}$ - 10^{-6}) were 1-2 orders of magnitude lower than estimates from observations (at the range of 10^{-4} - 10^{-5} for SO_4^{2-} , NO_3^- and NH_4^+ , respectively).*

L60-64: Here you argue that uncertainties in below-cloud scavenging coefficients is a potential reason for model underestimation of nitrate and sulfate wet deposition. How important are other sources of model uncertainties, such as N and S emissions, chemical transformation, changes in other ambient N and S compounds, etc.?

[Response]: Thanks for the comment. The other sources of model uncertainties, such as N and S emissions, chemical transformation, changes in other ambient N and S compounds, are also important to the simulation of nitrate and sulfate wet deposition. It is revised as: *This could be one reason for the underestimation of SO_4^{2-} and NO_3^- wet deposition in regional models of Asia reported in phase II and III of the Model Inter-Comparison Study for Asia (MICS-Asia) (Wang et al., 2008; Itahashi et al., 2020; Ge et al., 2020) and in global model assessments by the Task Force on Hemispheric Transport of Atmospheric Pollutants (TF-HTAP) (Vet et al., 2014), in addition to the other sources of model uncertainties (Chen et al., 2019; Tan et al., 2020; Kong et al., 2020), such as emissions, chemical transformation and changes in other ambient compounds of sulfur and nitrogen.*

L78-80: “The chemical components in later increments of rainfall are thought to be less influenced by the below-cloud scavenging process than by the in-cloud rainout process”. The wording is confusing. Why does the latter precipitation increments in sequential sampling represent in-cloud scavenging, while the start of the precipitation increments represent below-cloud scavenging? At what point during a precipitation event does the dominant wet scavenging process change from below-cloud to in-cloud? Is this transition point the same for all precipitation events and locations? Each precipitation event is unique in terms rainfall intensity, droplet sizes and distribution, precipitation form (rain, snow, or sleet), air concentrations of chemical components, etc. Meteorology varies with location; some locations are frequently affected by thunderstorms or deep convective scavenging. Can you address whether these factors are taken into consideration when using the sequential precipitation sampling method to estimate incloud or below-cloud scavenging contributions?

[Response]: Thanks for the comment. Similar as the response to the general comments

2 and 4. The initial concentration of raindrops in cloud is well mixed and can be considered as a stable state during the whole rainfall event. Actually, many observations in different regions (Aikawa *et al.*, 2009; 2014; Wang *et al.*, 2009; Quyang *et al.*, 2015; Xu *et al.*, 2017) reported that the chemical components in a rainfall event show a decayed variation with the increase of precipitation amount and eventually tends to a stable and low concentration level. The assumption in this study as well as the previous studies “*the concentrations in later increments can be attributed to scavenging by rainout only*” is based on this fact. It does not mean the below-cloud and in-cloud scavenging occur in sequence. But, instead, the two processes have been mixed in all stage of the rainfall event with the below-cloud scavenging contributed more in beginning fraction and the in-cloud scavenging contributed more in the later fraction due to the depletion of the air pollutants below cloud by washout.

The question is, how we recognized the “later fraction” in a rainfall event. The 5 mm accumulated precipitation (the concentration of chemical components at the 5th fraction) was used in the previous studies. However, in many cases, the concentration of chemical ions after 5 mm accumulated precipitation cannot be reached at stable level due to different precipitation intensity and the concentration level of air pollutants in each rainfall event. In this study, the asymptote value from the exponential decay fitting curve of the observed rainwater concentrations was employed as the in-cloud ion concentration. To investigate the uncertainties of this approach, the comparison between the exponential approach and the average value in fractions 6 to 8 (here after, average approach) has been carried out. The results show the exponential approach gives lower estimates of in-cloud concentrations than the average approach, with the latter being recognized as an upper limit for in-cloud concentrations.

And indeed, each precipitation event is unique in terms of rainfall intensity, rainfall type, air concentrations of chemical components, etc. The exponential approach to each unique event was made. In this sense, the unique meteorology conditions as well as air pollutions are considered in each rainfall event, since the variations of the ions concentration in each fraction of each rainfall are influenced by these synthetic effects. Besides, the estimation of below-cloud scavenging proportion based on the average value as well as the first quartile and the third quartile value are also included in the Figure 4 in addition to the median value to show a robust decreasing trend of below-cloud scavenging in wet deposition during 2014-2017. These have been revised in the new version.

L105: How much precipitation is collected in the eighth fraction? These details should be included in the paper.

[Response]: Agree. The eighth fractions in each rainfall event are different. The detail information has been added in the revised manuscript as: *During 2014-2017, a total of 104 precipitation events, which is almost 690 precipitation samples, were collected. Of the total number of precipitation events, 33 events (32%) were discarded from the sequential sampling analysis due to low rainfall amounts (<8 mm), which cannot satisfy the rules of full events. Altogether, 69 full events including 6 extended events were recorded over the 2014-2017 period in Beijing, as 15, 16, 20 and 18 events at each year,*

respectively. The rainfall volume of the eighth fraction of these 69 full events varied from 1 mm to 55.9 mm.

L109: In the 69 full events, what were the total precipitation amounts collected? Are they roughly the same for each event or do they vary greatly? At what point do you have to conduct manual sampling to collect the remainder of the precipitation? These details should be included in the paper.

[Response]: Agree. Similar as the response to major points 1 of the Anonymous Referee #1. During 2014-2017, a total of 104 precipitation events, which is almost 690 precipitation samples, were collected. Of the total number of precipitation events, 33 events (32%) were discarded from the sequential sampling analysis due to low rainfall amounts (<8 mm), which cannot satisfy the full events. The total precipitation amounts over 2014-2017 were 2151 mm with the data in each year listed in Table S1. The rainfall volume varied greatly for each event. Since mainly precipitation events occurred in night. The manual sampling has only been conducted in several heavy rainfall events which occurred in daytime. More details introduction on sampling have been added in the revised version as: *For example, if there is 12 mm rainfall volume in a precipitation event, 1 mm sequential rainfall is collected in each of the first 7 fractions with the rest of 5 mm in the eighth fraction. Rainfall events where eight fractions are collected and identified as full events, and those with fewer than eight fractions are characterized as incomplete events. Manual sampling methods were used to collect more than eight fractions during heavy rainfall, and these are characterized as extended events. During 2014-2017, a total of 104 precipitation events, which is almost 690 precipitation samples, were collected. Of the total number of precipitation events, 33 events (32%) were discarded from the sequential sampling analysis due to low rainfall amounts (<8 mm), which cannot satisfy the rules of full events. Altogether, 69 full events including 6 extended events were recorded over the 2014-2017 period in Beijing, as 15, 16, 20 and 18 events at each year, respectively. The rainfall volume of the eighth fraction of these 69 full events varied from 1 mm to 55.9 mm.*

L141-147: Is there additional evidence that can be presented to justify the assumption that precipitation at the start of an event represents below-cloud scavenging while latter precipitation represents in-cloud scavenging? Some studies have measured concentrations of chemical components in cloud water as an indication of in-cloud scavenging; however, the method in this study are based solely on the precipitation at the surface. One initial criticism I have on this assumption is why below-cloud and in-cloud processes have to occur one after another instead of simultaneously. A full justification is necessary given that the major results of this study depend on this analysis and interpretation.

[Response]: Thanks for the comment. According to the reported concentrations of chemical components in cloud water (Seinfeld and Pandis, 2006), which is 4-5 times higher than that observed at the surface. However, this can not be an indication of in-cloud scavenging, since the concentration might be diluted as the rainfall occurred. The direct observation just under the base of the cloud should be the concentration in-cloud.

Unfortunately, this measurement was absent in this study.

Similar as the response to the general comments 2. The assumption in this study does not mean the below-cloud and in-cloud scavenging occur in sequence. But, instead, the two processes have been mixed in all stage of the rainfall event with the below-cloud scavenging contributed more in beginning fraction and the in-cloud scavenging contributed more in the later fraction due to the depletion of the air pollutants below cloud by washout. The more detailed explanation on the estimation of below cloud scavenging including the full justification has been added in the revised manuscript.

L157: In previous studies, it seems that the 5 mm accumulated precipitation cut-off was based on the point where there was a lack of change in the precipitation concentrations. This is not entirely subjective.

[Response]: Agree. The expression has been revised as: *Previous studies have estimated the concentration of chemical ions scavenged in-cloud based on the judgment that 5 mm of accumulated precipitation is sufficient to identify the contribution of the rainout process (Wang et al., 2009; Aikawa and Hiraki, 2009; Xu et al., 2017).*

L164-174: Clarity on the methodology is needed. You mentioned that each precipitation event is unique in terms of the decreasing rate in the precipitation concentrations. What is the reason for combining all the precipitation events prior to fitting a regression curve? Why not fit a curve for each event? What is the purpose of fitting the data through the first and third quartiles – can this serve as an estimate of the uncertainties? “In theory, the concentration of chemical ions stabilize at higher rainfall increments and this represents the concentration in cloud.” A brief explanation of the theory should be provided in the paper.

[Response]: Agree. In this study, the fitting curve is implemented both in each rainfall event and in all combining events over 2014-2017. The latter is used to compare with the yearly scale in-cloud concentration which is reported in previous study from the median data. The first and third quartiles served as an estimate of the range of the uncertainties. The comparison between the exponential approach and the average approach (average value after 5th fraction, i.e., 6th-8th) has been carried out. The results show the exponential approach gives lower estimates of in-cloud concentrations than the average approach, with the latter being recognized as an upper limit for in-cloud concentrations. Based on this uncertainty analysis, the exponential approach to each unique event was made. This methodology has been revised to more clarity to the readers. It was now revised as: *In this study, a new method based on fitting a curve to the chemical ion concentrations with successive rainfall increments has been developed to estimate the contribution of the rainout process. As shown in Figure 1, an exponential curve is fitted to the median, 25th and 75th percentiles of the chemical ion concentrations in each fraction through the rainfall increments. Noted that, the fitted exponential curve is applied to the combination of all 69 full events to estimate the yearly median concentration of chemical ions in-cloud and to compare with the results from previously reported method (i.e., median concentration after 5 mm increments). Besides, the exponential approach to each unique event was also employed. Ideally, the*

concentration of chemical ions stabilize at higher rainfall increments and this represents the concentration in cloud. However, the decrease during each rainfall event is distinctly different, and this regression method is not fully applicable to all rainfall events in practice. Therefore, the exponential regression method is used to estimate the in-cloud concentration under most circumstances, but where the decreasing trend with the increment of rainfall is not significant, the average value of rainfall increments 6-8 of the event is used.

Equations 1,2: What does n represent? n in eq. 1 is different from the n in eq. 2.

[Response]: n represent the total fractions in a rainfall event. This has been added in the revised manuscript.

L185: Some background information on the Action Plan is needed in the introduction to understand what effects these policies may have on air quality. Which pollutants does the Action Plan target and what was implemented?

[Response]: The Action Plan is launched in 2013 called “Ten rules” to improve the air quality in China. It includes comprehensive control of industrial emission, non-point emission, fugitive dust, vehicles, etc. It is also aimed to adjust and optimize the industrial structure and promote economic transformation and upgrading, such as increase the supply of clean energy. These actions are ensured to work by both of legislation and market mechanism. According to the *Beijing Environmental Statement* published by the Beijing Municipal Environmental Protection Bureau from 2013 to 2017, many measures have been implemented to meet the Action Plan, including replacement residential coal with electricity and natural gas, upgrading the emission standards of gasoline, diesel vehicles and power plants, closing the high-emission enterprises. This description has been added in section 3.1 of the revised manuscript.

Section 3.1: Why are the results of other inorganic ions not discussed? The annual changes in the VWM concentrations should be discussed with changes in emissions of the precursor pollutants, such as SO_x and NO_x emissions, and policies that were introduced to manage those emissions. There should also be more discussion on what was causing the significant declines in sulfate and Ca²⁺ in the earlier period. Air pollution control measures typically target major anthropogenic contaminants, but here you have shown significant decreases in Ca²⁺ which is largely from soil emissions. It is not clear how the Action Plan played a role in the reduction of VWM Ca²⁺. For a better understanding of the impacts of acidification on ecosystems, wet deposition fluxes should be presented as well. While there may be significant decreases in VWM concentrations, interannual variability in precipitation amounts may result in less significant decreases in wet deposition fluxes.

[Response]: In this study, the major ions in precipitation are considered. According to the measurement, the two anions (SO₄²⁻ and NO₃⁻) and two cations (NH₄⁺ and Ca²⁺) are discussed in the main text with the other ions listed in Table 2 and plotted in Figure 7 in the revised manuscript.

The annual changes in VWM concentrations with changes in emissions and the

annual concentrations of SO₂ and NO_x have been added in section 3.1 of the revised manuscript. The emission and the concentration data of SO₂ and NO_x are collected from the yearly book of “*Environmental Bulletin in Beijing*” from 1994 to 2017. It is clearly shown the concentration of SO₂ experienced a sustainably decreasing trend due to significant reduction of its emission from 1996 to 2017, with the decreases rate is 4.5% yr⁻¹ and 13.9% yr⁻¹ in emission and 2.8% yr⁻¹ and 14.0% yr⁻¹ in concentration during 1995-2010 and 2013-2017 (the Action Plan period), respectively. This is consistent with the annual changes of VWM concentrations of SO₄²⁻. As to NO_x emission, the data in recent years have been collected. Although the clearly reduction is found in the annual changes of emission from 2010, the ambient concentration of NO₂ do not show a significant decreasing trend (~3.6% yr⁻¹) compared with SO₂ (14% yr⁻¹). However, before the Action Plan, the decreasing ratio in concentration is only 1.8% yr⁻¹, which is slower than the Action Plan period. Different from its variation in the ambient concentration, the increase of VWM concentration of NO₃⁻ (+4% yr⁻¹) in precipitation is found during 1995-2010 and then decrease (-3% yr⁻¹) in 2013-2017. Both of the annual changes in SO₂ and NO₂ implies the dominant contribution of the Action Plan to the air quality.

The significant declines in VWM concentration of Ca²⁺ is found in precipitation with the decreases rate as 36.1% yr⁻¹ in 1995-2010 and 8.8% yr⁻¹ in 2014-2017. The emission and the concentration data of Ca²⁺ are absent in this study. Instead, the different of PM₁₀ and PM_{2.5} (PM₁₀-PM_{2.5}) air concentration during 2013-2017 have been calculated to represent the coarse particles, which contains the Ca²⁺ and Na⁺ as well. The results show that the concentration decreased from 31.2 µg/m³ in 2013-2014 to 24.0 µg/m³ over 2015-2017. This indicates the improvement of coarse particles even which is derived from crustal emissions have been made through the Action Plan. Similar as the response to “specific comments L185”, the Action Plan launched in 2013 including emission reduction not only from energy consumption of industry but also the fugitive dust in cities, which should result the decline in Ca²⁺.

Observations on S and N wet deposition (Pan et al., 2012; 2013) during 2007-2010 show the value of 21.5 kg S ha⁻¹ yr⁻¹ and 27.9 kg N ha⁻¹ yr⁻¹ (19.7 and 8.2 kg N ha⁻¹ yr⁻¹ through NO₃⁻ and NH₄⁺) in Beijing, respectively. Compared with those results, significant decreases (11.4 kg S ha⁻¹ yr⁻¹ and 23.6 kg N ha⁻¹ yr⁻¹) were observed in the four-years measurements during 2014-2017 in this study.

Further analysis on annual changes listed above have been added in section 3.1 of the revised manuscript.

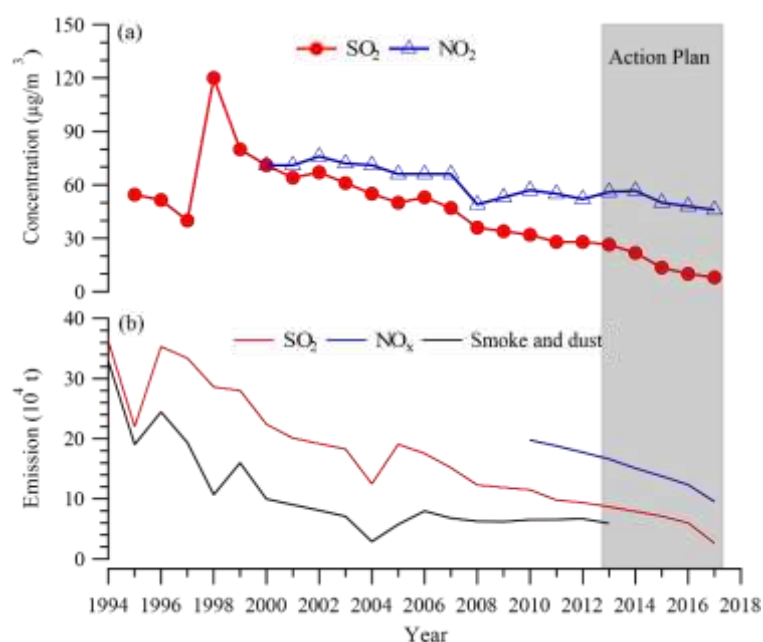


Figure 3 (in the revised manuscript). Annual changes in emission and concentration of SO₂ and NO_x in Beijing (collected from the yearly book of “*Environmental Bulletin in Beijing*” from 1994 to 2017).

L219: *H* usually denotes a Henry’s Law constant. It is more suitable to use *W* or *SR* to denote scavenging ratios as this is consistent with literature.

[Response]: Agree. The *H* has been replaced by *W* in the revised manuscript.

L234: “while the *VWA* represents a greater contribution from in-cloud removal: :” It was stated earlier that latter fractions of the accumulated precipitation represents incloud removal. Why was the *VWA* concentrations (which may include below-cloud and in-cloud contributions) used instead of the concentrations in the latter precipitation fractions?

[Response]: It is the *VWA* concentrations may include below-cloud and in-cloud contributions. The relations between the concentration of ions in *VWM* and the ambient air shows the synthetic correlations. Based on the comparison of the relations between air and precipitation from the first fraction or from *VWM*, we can get not only the influences from the air pollutants below-cloud but also the uncertainties due to coarse particles and gases below-cloud (especially to HNO₃ and NH₃). The larger difference of the *R* coefficients for NO₃⁻ and NH₄⁺ than SO₄²⁻ in Figure 4 (original Figure 3) may indicate the uncertainties. The former two ions are related to more complicate sources from the ambient precursors. For example, the NO₃⁻ in precipitation is both from the fine and coarse particles (i.e., particulate NO₃⁻) as well as the gaseous HNO₃, while the NH₄⁺ in precipitation is mainly from the fine particles in addition to NH₃.

L237-238: “This indicates that the concentration of chemical ions in precipitation at the start of rainfall is more greatly influenced by aerosols below the cloud.” This conclusion is not correct because the scavenging of acidic gases like HNO₃ and SO₂ can also contribute to nitrate and sulfate in precipitation. Another issue with the correlation analysis between precipitation and air concentrations (in Fig. 3) is that only

fine aerosols were considered. Would there be any changes to the correlation results if acidic gases and coarse aerosols were included?

[Response]: Similar response to general comment 1. To avoid misunderstanding, this sentence has been revised as: *This indicates that the concentration of chemical ions in precipitation at the start of rainfall is more greatly influenced by the air pollutants below the cloud.*

L245: The scavenging ratios of SNA do not include coarse particles since only PM_{2.5} chemical composition were measured. It also does not include the wet scavenging of acidic gases. This needs to be mentioned in the paper because typically scavenging ratios take into account all the chemical species in air that can undergo wet deposition. It seems that the scavenging ratios of SNA calculated in this study is not an accurate reflection of the wet scavenging efficiency of SNA.

[Response]: Agree. The scavenging ratios of SNA in this study is not an accurate reflection of the wet scavenging efficiency of SNA due to lacking of consideration of coarse particles and acidic gases. Similar as the response to general comment 1, the absent measurement to coarse particles and acidic gases may lead to little influence to SO_4^{2-} , but large uncertainties in NO_3^- and NH_4^+ . To avoid misunderstanding, this has been clarified as: *It should be noted that the W calculated in this study is based on the fine particles in air, which may not represent the accurate reflection of the wet scavenging efficiency of SNA.*

L264-266: The percentages should be converted to annualized percentages. Some of the time periods are much longer than others, which makes it appear that the rate of decrease is larger.

[Response]: Agree. The percentages here have been revised to the annualized percentages in the revised manuscript.

L269: "The chemical composition of precipitation is directly related to the amount of precipitation, and as a consequence it is difficult to identify inter-annual variations in chemical concentrations." This statement is incorrect. There are other factors affecting the precipitation chemistry besides the precipitation amount. Section 3.3: An exponential curve was used to fit the data points and then the horizontal asymptote was used to determine the precipitation concentration representative of rainout. One issue that I have with this approach is that there were only 11 data points or precipitation fractions. Previous studies (e.g. Aikawa and Hiraki, 2009) that conducted sequential sampling of precipitation found that the precipitation concentrations could increase in the much latter fractions of precipitation. It seems that more precipitation fractions need to be collected in order to observe what happens to the precipitation concentrations.

[Response]: Agree. There are many other factors that affecting the precipitation chemistry. In this study, we also observed the increased precipitation concentrations in the latter fractions of precipitation. This may due to the meteorological conditions, i.e., rainfall type, intensity, cloud and the air pollution movement. Previous studies (e.g.

Aikawa and Hiraki, 2009) that conducted sequential sampling of precipitation were based on 0.5 mm increment and found the low level concentration remained (in-cloud concentration) at after 10 fractions, which is 5 mm accumulated rainfall. In this study, the sequential sampling of 1 mm increment with 8 fractions at least is employed, which covering higher than 5 mm accumulated rainfall. However, despite the longer precipitation fractions in this study were collected, the longer fraction measurement and more detailed analysis on the uncertainties are needed in the future. In this study, we agreed with the reviewer's comment and deleted this sentence in the revised manuscript.

L303-308: "Benefiting from the Action Plan, the below-cloud contributions: : :" This part may not be necessary because it was discussed earlier that the policies in the Action Plan help reduced the VWM of inorganic ions. The Action Plan did not directly affect the below-cloud scavenging process. Rather, it improved the air quality which in turn decreased the VWM concentrations.

[Response]: Thanks for the comment. We still insist to keep this part in the revised manuscript based on the following reasons:

- 1) Yes. It was discussed earlier that the policies in the Action Plan help reduced the VWM of inorganic ions. However, the VWM of inorganic ions in precipitation has two part (i.e., in-cloud and below-cloud contribution) in general. This part gives more detailed information that the decreases of VWM may largely due to the decreases of below-cloud part.
- 2) We agree with that the Action Plan did not directly affect the below-cloud scavenging process. However, the air concentration at the surface layer have been significant reduced under the implementation of the Action Plan. This improved the air quality at the surface layer and in turn decreased the below-cloud scavenging in the total wet deposition.

To avoid misunderstanding, the expression here has been revised as: *Benefiting from the Action Plan, the air quality at the surface layer have been significantly improved (Zhang et al., 2019), which in turn leading to the decreases of the below-cloud scavenging. In this study, it also shows the below-cloud contributions of SO_4^{2-} , NO_3^- , NH_4^+ and Ca^{2+} decreases from >50% in 2014 to ~40% in 2017.*

Section 3.4: I think this section is not necessary because the findings are not substantially new from what has already been discussed in the paper. Here you are showing the effects of rainfall on ambient air sulfate. The washout effects were already discussed in Figure 1, which shows the decrease in VWM concentrations of the inorganic ions with increasing rainfall.

[Response]: Agree. The section 3.4 has been removed from the revised manuscript.

L392: "highest contribution for NH_4^+ at 65% and lowest for SO_4^{2-} at 50%". This was stated in the conclusion and abstract, but was not elaborated in the results. I expected the highest contribution to below-cloud wet scavenging would have been from Ca^{2+} or Na^+ given that they are predominantly associated with coarse particles and locally emitted.

[Response]: Agree. Considering the uncertainties of below-cloud scavenging contribution to NH_4^+ wet deposition, the expression has been deleted both in abstract and in conclusion. Besides, it is interesting to note that the yearly below-cloud contribution to total wet deposition of Ca^+ decreased from >60% to less 40% during 2014-2017. This may due to the clean air Action Plan, which sustainably decreased the coarse particles with the different of PM_{10} and $\text{PM}_{2.5}$ ($\text{PM}_{10}-\text{PM}_{2.5}$) decreased from $31.2 \mu\text{g}/\text{m}^3$ in 2013-2014 to $24.0 \mu\text{g}/\text{m}^3$ over 2015-2017. Similar as the response to “specific comments L185 and section 3.1”, the Action Plan launched in 2013 including emission reduction not only from energy consumption of industry but also the fugitive dust in cities, which should result the decline in Ca^{2+} .

References:

- Aikawa, M., and Hiraki, T.: Washout/rainout contribution in wet deposition estimated by 0.5 mm precipitation sampling/analysis, *Atmos Environ*, 43, 4935-4939, 2009.
- Aikawa, M., Kajino, M., Hiraki, T., and Mukai, H.: The contribution of site to washout and rainout: Precipitation chemistry based on sample analysis from 0.5 mm precipitation increments and numerical simulation, *Atmos Environ*, 95, 165-174, <http://dx.doi.org/10.1016/j.atmosenv.2014.06.015>, 2014.
- Chen, L., Gao, Y., Zhang, M., Fu, J. S., Zhu, J., Liao, H., Li, J., Huang, K., Ge, B., Wang, X., Lam, Y. F., Lin, C. Y., Itahashi, S., Nagashima, T., Kajino, M., Yamaji, K., Wang, Z., and Kurokawa, J.: MICS-Asia III: multi-model comparison and evaluation of aerosol over East Asia, *Atmos. Chem. Phys.*, 19, 11911-11937,
- Ge, B., Itahashi, S., Sato, K., Xu, D., Wang, J., Fan, F., Tan, Q., Fu, J. S., Wang, X., Yamaji, K., Nagashima, T., Li, J., Kajino, M., Liao, H., Zhang, M., Wang, Z., Li, M., Woo, J. H., Kurokawa, J., Pan, Y., Wu, Q., Liu, X., and Wang, Z.: MICS-Asia III: Multi-model comparison of reactive Nitrogen deposition over China, *Atmos. Chem. Phys. Discuss.*, 2020, 1-43, [10.5194/acp-2019-1083](https://doi.org/10.5194/acp-2019-1083), 2020.
- Itahashi, S., Ge, B., Sato, K., Fu, J. S., Wang, X., Yamaji, K., Nagashima, T., Li, J., Kajino, M., Liao, H., Zhang, M., Wang, Z., Li, M., Kurokawa, J., Carmichael, G. R., and Wang, Z.: MICS-Asia III: overview of model intercomparison and evaluation of acid deposition over Asia, *Atmos. Chem. Phys.*, 20, 2667-2693, [10.5194/acp-20-2667-2020](https://doi.org/10.5194/acp-20-2667-2020), 2020.
- Kajino, M., and Aikawa, M.: A model validation study of the washout/rainout contribution of sulfate and nitrate in wet deposition compared with precipitation chemistry data in Japan, *Atmos Environ*, 117, 124-134, [10.1016/j.atmosenv.2015.06.042](https://doi.org/10.1016/j.atmosenv.2015.06.042), 2015.
- Kasper-Giebl, A., Kalina, M. F., and Puxbaum, H.: Scavenging ratios for sulfate, ammonium and nitrate determined at Mt. Sonnblick (3106 m asl), *Atmos Environ*, 33, 895-906, 1999.
- Kong, L., Tang, X., Zhu, J., Wang, Z., Fu, J. S., Wang, X., Itahashi, S., Yamaji, K., Nagashima, T., Lee, H. J., Kim, C. H., Lin, C. Y., Chen, L., Zhang, M., Tao, Z., Li, J., Kajino, M., Liao, H., Wang, Z., Sudo, K., Wang, Y., Pan, Y., Tang, G., Li, M., Wu, Q., Ge, B., and Carmichael, G. R.: Evaluation and uncertainty investigation of the NO_2 , CO and NH_3 modeling over China under the framework of MICS-

- Asia III, *Atmos. Chem. Phys.*, 20, 181-202, 10.5194/acp-20-181-2020, 2020.
- Ouyang, W., Guo, B. B., Cai, G. Q., Li, Q., Han, S., Liu, B., and Liu, X. G.: The washing effect of precipitation on particulate matter and the pollution dynamics of rainwater in downtown Beijing, *Sci Total Environ*, 505, 306-314, 2015.
- Seinfeld, J. H., and Pandis, S. N.: *Atmospheric chemistry and physics: from air pollution to climate change*, Wiley, New York, 2006.
- Tan, J., Fu, J. S., Carmichael, G. R., Itahashi, S., Tao, Z., Huang, K., Dong, X., Yamaji, K., Nagashima, T., Wang, X., Liu, Y., Lee, H. J., Lin, C. Y., Ge, B., Kajino, M., Zhu, J., Zhang, M., Liao, H., and Wang, Z.: Why do models perform differently on particulate matter over East Asia? A multi-model intercomparison study for MICS-Asia III, *Atmos. Chem. Phys.*, 20, 7393-7410, 10.5194/acp-20-7393-2020, 2020.
- Vet, R., Artz, R. S., Carou, S., Shaw, M., Ro, C. U., Aas, W., Baker, A., Bowersox, V. C., Dentener, F., Galy-Lacaux, C., Hou, A., Pienaar, J. J., Gillett, R., Forti, M. C., Gromov, S., Hara, H., Khodzher, T., Mahowald, N. M., Nickovic, S., Rao, P. S. P., and Reid, N. W.: A global assessment of precipitation chemistry and deposition of sulfur, nitrogen, sea salt, base cations, organic acids, acidity and pH, and phosphorus, *Atmos Environ*, 93, 3-100, 2014.
- Wang, Y., Xue, L. I., Li, Y. A. O., Yanan, Z., and Yuepeng, P. A. N.: Variation of pH and Chemical Composition of Precipitation by Multi-step Sampling in Summer of Beijing 2007 (in english abstract), 30, 2715-2721, 2009.
- Xu, D. H., Ge, B. Z., Wang, Z. F., Sun, Y. L., Chen, Y., Ji, D. S., Yang, T., Ma, Z. Q., Cheng, N. L., Hao, J. Q., and Yao, X. F.: Below-cloud wet scavenging of soluble inorganic ions by rain in Beijing during the summer of 2014, *Environ Pollut*, 230, 963-973, 10.1016/j.envpol.2017.07.033, 2017.
- Yue, D., Hu, M., and Wu, Z.: Variation and interaction of major azotic inorganic compounds in the summer in beijing (in Chinese), *Environ. Monit. China*, 29, 9-14, 2013.
- Wang, Z. F., Xie, F. Y., Sakurai, T., Ueda, H., Han, Z. W., Carmichael, G. R., Streets, D., Engardt, M., Holloway, T., Hayami, H., Kajino, M., Thongboonchoo, N., Bennet, C., Park, S. U., Fung, C., Chang, A., Sartelet, K., and Amann, M.: MICS-Asia II: Model inter-comparison and evaluation of acid deposition, *Atmos Environ*, 42, 3528-3542, 2008.