

Interactive comment on “Limited production of sulfate and nitrate on front-associated dust storm particles moving from desert to distant populated areas in northwestern China” by Feng Wu et al.

Anonymous Referee #2

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General comments:

A number of laboratory and field studies have proved that Asian dust particles readily promoted the formation of sulfate and nitrate when the lofted dust plumes transported across urban areas under high RH and elevated levels of reactive trace gases (i.e. SO₂, NO_x, O₃ and OH radicals). This would significantly alter the physical and chemical properties of dust aerosol and subsequent climate change on regional scale.

The authors carried out a series of particle samplings at the Tengger desert (06:30–15:00 BST on April 24, 2014) and downwind Xi’an city (07:00–19:00 BST on May 1, 2014) during two independent dust storms. Combination of HYSPLIT backward trajectories model and CFORS simulation, they showed that the two dust events originated

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from the same source regions and had similar transport routes. They compared the concentrations and mass fraction of chemical components (i.e. sulfate, nitrate, ammonia, and elemental ratios) in dust particles at two sites during the prefrontal, frontal, and postfrontal air parcels, and indicated that the production of sulfate and nitrate on front-associated dust particles was limited when the dust moved from desert sources to populated areas in northwest China. The result of this manuscript seems to be reasonable in spite of limited in-situ sampling data, which is completely different from the other previous studies. Different scientific viewpoints should be encouraged to promote the understanding of interplay between mineral dust and atmospheric chemistry. Therefore, I recommend this manuscript is accepted and published in the journal of ACP after some revisions.

Specific comments:

1. Abstract, Page 2, lines 12–14: “The significant sulfate and nitrate reported in dust-associated samples in previous studies were more likely produced on locally-emitted and urban mineral particles or from soil-derived sulfate rather than being produced via chemical conversions on desert dust particles.”

Conclusion, Page 11, lines 2–4: “Significant sulfate and nitrate in dust storm periods in China reported in previous studies were likely produced on locally-emitted and urban mineral particles, in addition to soil-derived sulfate, and they were unlikely produced via chemical conversions on dust particles from deserts.”

Comment: I think there is no enough evidence for the manuscript to demonstrate this conclusion. Because the dry condition ($RH < 40$)

2. Page 4, lines 21–22: “This sample collection ensured that mineral particles collected on the filters were dust particles from the desert and there should be no influence of anthropogenic pollutants from the village or the city considered in the samples.”

Comment: The evidence provided by this manuscript could not fully support this sen-

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tence. Please reconsider again.

3. Page 6, lines 15–17: “The cold fronts are the boundaries between the local or regional anthropogenic-polluted air and the long-distance transported air because the movement of air on a synoptic scale is approximately adiabatic, i.e. the air is hardly mixed with thermodynamically-different air it meets.”

Comment: I don't agree with this viewpoint about “the air is hardly mixed with thermodynamically-different air it meets”. In terms of meteorology, the warm and humid air mass is readily lifted and the weather process (e.g., strong wind and cooling weather, rainfall or snow) often changes dramatically on the border of frontal system when a cold front passes over. As shown in Figure 2a, the RH increased sharply from 40

4. Page 7, lines 26–33: “At the desert site, NO₃- concentration in dust samples was 4-6 $\mu\text{g m}^{-3}$ and the average was 5 $\mu\text{g m}^{-3}$. The relative amount of NO₃- range between 0.11

Comment: At the Tengger desert site, NO₃- concentration in dust samples was 4-6 $\mu\text{g m}^{-3}$ (with the average value and fraction of 5 $\mu\text{g m}^{-3}$ and 0.12

5. Page 18, Table 2; Page 19, Table 3: The authors sampled the concentrations of TSP (total suspended particulates) and analyzed the chemical components (i.e. sulfate, nitrate, and ammonia) in TSP at the Tengger desert and Xi'an sites.

Comment: Please explain why did you sample the concentrations of TSP, instead of PM₁₀ or PM_{2.5}. It is well known that most of the coarse-size dust particles (radii > 10 μm) generally settle near the source region on account of large gravitational deposition velocity, whereas the finer dust particles (radii < 10 μm) are transported more efficiently to the downstream areas. And the concentrations of TSP (meaning coarse-size particle with radii > 10 μm) in Xi'an city should include the local source emissions (e.g., engines of vehicles, road dust, and construction dust; Page 4, lines 27-29) that increases the

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TSP concentrations, but may decrease the relative mass fraction of sulfate, nitrate, and ammonia. Inferred from Page 6, Lines 1-7, the mass concentrations of TSP and sulfate are about 425 $\mu\text{g m}^{-3}$ and 17 $\mu\text{g m}^{-3}$ at Xi'an before the dust arrival (i.e. prefrontal air). In the postfrontal air, the mass concentrations of sulfate are 3.8, 3.5, and 3.4 $\mu\text{g m}^{-3}$, respectively, right after, two hour after, and four hours after the passage of cold front (means slight variations), whereas the corresponding TSP concentrations are 422, 318, and 189 $\mu\text{g m}^{-3}$ (shows large variations). Clearly, relative mass fractions of sulfate reduce.

Minor comments:

1. Abstract, Page 2, lines 3–4: “but the production was very inefficient in other studies.”

Comment: Please give the quoted literature.

2. Page 3, line 8: “RH”

Comment: Change to “relative humidity (RH)”

3. Page 10, line 4: “mineral/TSP ratios”

Comment: Change to “mineral/TSP (total suspended particulates) ratios”

Please also note the supplement to this comment:

<http://www.atmos-chem-phys-discuss.net/acp-2016-853/acp-2016-853-RC2-supplement.pdf>

Interactive comment on Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-853, 2016.

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