1 Dear editor,

Thank you for your kind considerations on our manuscript entitled "Aerosol-2 meteorology feedback diminishes the trans-boundary transport of black carbon into the 3 Tibetan Plateau" (egusphere-2023-252). We appreciate that you gave us a chance to 4 improve our manuscript to a level suitable for publication in ACP. We also want to 5 6 express our deep thanks to the reviewers of the positive comments. Those comments are all valuable and very helpful for revising and improving our paper. We have studied 7 comments carefully and have made corrections, which we hope meet with approval. 8 The main corrections in the paper and the responds to the reviewer's comments are as 9 following: 10

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12 Answers to reviewers:

13 Reviewer #2:

The interaction between aerosols and meteorology, and its impact on the cross-14 boundary transport flux of BC (black carbon) over the Tibetan Plateau (TP), has 15 received limited attention in previous research. This paper presents a comprehensive 16 investigation of the aerosol-meteorology feedback and its influence on BC transport 17 flux during a period of heavy aerosol pollution. The study utilizes WRF-Chem 18 19 simulation to thoroughly analyze the phenomenon. Additionally, the paper elucidates 20 the meteorological factors that contribute to the occurrence of severe aerosol pollution events over the TP. The concept introduced in this article is characterized by its novelty, 21 and the study's findings hold significant implications for the preservation of the TP's 22 ecological environment. Hence, I recommend that this manuscript be revised and 23 24 considered for publication in ACP. Please find below some specific comments for 25 further improvement:

Firstly, we appreciate that you gave us a chance of revision to improve our manuscript to a level suitable for publication in Atmospheric Chemistry and physics. The comments are replied as follows:

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The authors validated the model performance on BC and AOD by comparing the
 simulation and observation. Although the comparison results are basically satisfactory,
 the data used to validate the model performance is still simple and I suggest inter-model
 comparison should be considered, which might be more convincing.

Response: Thank you very much for your valuable advice. The reviewer made a very 34 good point here. According to the reviewer's suggestion, we not only validated the 35 model performance on temporal variation in AOD at different stations by comparing 36 37 the simulated AOD with the ground-based and satellite-based observational AOD (Figure S4), but also verified the model performance on the spatial distribution of AOD 38 over the study area by comparing the simulated AOD with the satellite-based and 39 reanalyzed AOD (Figure S5). However, for BC, the comparison between reanalysis and 40 simulation was only conducted because we have very limited in-situ observed BC (the 41 observed BC is only available at the QOMS station). According to the reviewer's 42 suggestion, we further validated the model performance by conducting the inter-43 comparison among in-situ observation, simulation, and MERRA-2 reanalysis, as shown 44

in Figure A1. The results show that the temporal variation in simulation is very close 45 to that of simulation. Moreover, the correlation coefficient between the simulation and 46 observation is 0.867, passing the 99% confidence level. Therefore, the model 47 configuration used in this study presented a reasonable performance on BC. For the 48 49 spatial distribution of BC over the study area, we compared the simulation with 50 reanalysis from MERRA-2 and simulation from CAM Chem. The results show that the spatial pattern of the WRF-Chem simulated BC is similar to that of the reanalyzed BC 51 (Figure S6); however, the spatial pattern of BC from CAM Chem is not reasonable 52 53 (Figure A2).

Additionally, in terms of the BC transport flux, as the spatial pattern of BC from CAM_Chem is not reasonable, we can't further verify the transboundary transport flux of BC with CAM_Chem data. Also, the BC from MERRA-2 has no vertical information, resulting in the inability to provide vertical profile of BC transport flux. Therefore, the transport flux of BC was not verified by inter-model comparison.



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60 Figure S4. Inter-comparison of temporal variations in simulated AOD and ground-61 based as well as satellite-based AOD at (a) Nam Co, (b) QOMS, and (c) Pokhara 62 stations for the period from April 20 to May 10, 2016

62 stations for the period from April 20 to May 10, 2016.

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Figure S5. Inter-comparison of spatial distribution of simulated mean daily AOD and
satellite-based as well as reanalyzed mean daily AOD from April 20 to May 10, 2016,
over the study area.

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71 Figure A1. Inter-comparison of temporal variations in simulated BC and in-situ

- observed BC as well as reanalyzed BC at QOMS station for the period from April 20
- 73 to May 10, 2016.
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Figure S6. Spatial distributions of simulated and reanalyzed daily mean BC
 concentrations over the domain averaged for the period from April 20 to May 10, 2016.

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80 Figure A2. Spatial distribution of daily mean BC from CAM_Chem data averaged for

81 the period from April 20 to May 10, 2016.

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83 2. When analyzing the meteorological causes of the heavy aerosol pollution event,
84 isotherms in the weather maps in Figure 3 are not included in the analysis, and isotherms

- lead to blurring of potential heights and wind fields in weather maps, so I suggest
- 86 removing them.
- 87 Response: Thank you very much for your kind remind. According to the reviewer's
- suggestion, we have removed the isotherms in the weather maps in Figure 3 and the
- 89 replotted Figure 3 is shown as follows:





Figure 3 Weather maps at 500 hPa over the study area during the severe aerosol pollution event based on ERA-Interim reanalysis dataset. The blue lines are isopleths of geopotential height (unit: dagpm). Wind speed (unit: m/s) and direction are denoted by wind barb.

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3. As the author stated in the title as well as in Figure.12, the aerosol-meteorology
feedback decreased the cross-boundary transport flux of BC towards the TP. In fact,
this conclusion is the result of pure model simulation because of the harsh environment,
limited access for fieldwork, and the sparsity of fixed instrumental stations over the TP.
So is there similar study in other places and What effect does the aerosol-meteorology
feedback have on the transport flux of aerosols?

Response: Thank you very much for your good suggestion. This suggestion is 102 somewhat similar to that made by the first reviewer. By reviewing extensive literature, 103 it was found that there are similar studies in other places. For example, using ground-104 based multi-axis differential optical absorption spectroscopy (MAX-DOAS) at the 105 Nancheng site in suburban Beijing on the southwest transport pathway of the Beijing-106 107 Tianjin-Hebei (BTH) region, Hu et at. (2022) estimated the vertical profiles of transport 108 fluxes in the southwest-northeast direction. The results showed that the maximum net 109 transport fluxes per unit cross-sectional area, calculated as pollutant concentration multiply by wind speed, of aerosol extinction coefficient (AEC), NO₂, SO₂ and HCHO 110 were 0.98 km⁻¹m s⁻¹, 24, 14 and 8.0 μ g m⁻² s⁻¹ from southwest to northeast, which 111 occurred in the 200-300 m, 100-200 m, 500-600 m and 500-600 m layers, respectively, 112 due to much higher pollutant concentrations during southwest transport than during 113 northeast transport in these layers. The average net column transport fluxes were 1200 114 $km^{-1} m^2 s^{-1}$, 38, 26 and 15 mg m⁻¹ s⁻¹ from southwest to northeast for AEC, NO₂, SO₂ 115

and HCHO, respectively, in which the fluxes in the surface layer (0–100 m) accounted
for only 2.3%–4.2%.

However, in terms of the influence of aerosol-meteorology feedback on transport 118 flux of aerosols, it is found that no matter in regions with abundant observational data 119 or in regions with sparse observational data, the influence of aerosol-meteorology 120 121 feedback on the transport flux of aerosols was evaluated by means of model simulation, because sensitivity tests are involved in such studies. For instance, Huang et al. (2020) 122 reported that long-range transport and aerosol-meteorology feedback may interact 123 rather than act as two isolated processes as traditionally thought by investigating typical 124 regional haze events in northern and eastern China. This interaction can then amplify 125 transboundary air pollution transport over a distance of 1,000 km and boost long-lasting 126 127 secondary haze from the North China Plain to the Yangtze River delta. The results show an amplified transboundary transport of haze by aerosol-meteorology interaction in 128 China and suggest the importance of coordinated cross-regional emission reduction 129 with a focus on radiatively active species like black carbon. 130

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4. Line 949, Line 1090, Line 1176, Line 1213–1214, and Line 1227: ATMOSPHERIC
CHEMISTRY AND PHYSICS --> Atmos. Chem. Phys.

- Response: Thank you for your suggestion. We are very sorry for our carelessness, and
 "ATMOSPHERIC CHEMISTRY AND PHYSICS" has been revised as "Atmos. Chem.
 Phys".
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138 5. Line 1128: JOURNAL OF GEOPHYSICAL RESEARCH-ATMOSPHERES -->
 139 Journal of Geophysical Research: Atmospheres

140 Response: Thank you for your suggestion. We have revised "JOURNAL OF
141 GEOPHYSICAL RESEARCH-ATMOSPHERES" as "Journal of Geophysical
142 Research: Atmospheres".

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144 6. Line 1034–1035: NATURE CLIMATE CHANGE -->Nature Climate Change

- 145 Response: Thank you for your suggestion. We have revised "NATURE CLIMATE146 CHANGE" as "Nature Climate Change".
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148 7. Line 1220: SCIENTIFIC REPORT--> Scientific Report

149 Response: Thank you for your advice. We have revised "SCIENTIFIC REPORT" as

- 150 "Scientific Report".
- 151

152 8. Line 1221–1224: The corresponding article is quoted incorrectly and lacks the 153 journal name.

154 Response: Thank you for your suggestion. The journal name has been added and the

155 correct citation is 'Zheng, B., Zhang, Q., Zhang, Y., He, K. B., Wang, K., Zheng, G. J.,

156 Duan, F. K., Ma, Y. L., and Kimoto, T.: Heterogeneous chemistry: a mechanism

157 missing in current models to explain secondary inorganic aerosol formation during the

158 January 2013 haze episode in North China, Atmos. Chem. Phys., 15 (4), 2031-2049,

159 https://doi.org/10.5194/acp-15-2031-2015, 2015.'.

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161	Once again, special thanks to you for your good comments.
162	Best Regards.
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164	Yuling Hu and Shichang Kang on behalf of all co-authors.
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166	Hu, Q., Liu, C., Li, Q., Liu, T., Ji, X., Zhu, Y., Xing, C., Liu, H., Tan, W., and Gao, M.: Vertical profiles of
167	the transport fluxes of aerosol and its precursors between Beijing and its southwest cities, Environ.
168	Pollut., 312, 119988, https://doi.org/https://doi.org/10.1016/j.envpol.2022.119988, 2022.
169	