

Response to Reviewer I

General Comments:

- Understanding the processes controlling the O₃ concentration in a specific area is important to design emission reduction strategies to reduce the harmful effects of tropospheric O₃. This paper focuses on two processes that take place in the O₃ cycle: transport and photochemistry.
- The paper discusses two methodological approaches to understand two O₃ processes (transport and photochemistry) which are the O₃ budget and the O₃ source apportionment. Authors claims that there is a contradictory view on the role of transport and photochemistry in O₃ pollution between the budget calculation studies and O₃ source apportionment studies, because both studies provide different information. In my point of view, they are two different approaches difficult to compare, so it is normal they provide different results. However, as the authors show in the paper it is possible to learn from the both of them.
- I think the paper is organized in a way that it does not help to understand its objective and methodology, even it shows a hard work behind. So, in my opinion, this manuscript was hard to follow and understand, and consequently to review. Furthermore, it could help if authors improve the readability of the text. Overall, there are too many pronouns and missing nouns that make difficult to follow the main idea of some sentences. Authors should review the text carefully and provide a more accurate reference to key concepts, also being consistent in the way they do it along the manuscript.

Response:

We appreciate the valuable comments and suggestions. We've tried to adjust the structure of the paper and make a lot of revisions to improve its readability.

Before in-detail responses, we want to clarify the “contradictory” in this paper. Reported O₃ concentration budgets often show that photochemistry is the main process leading to the rapid increase of O₃ concentrations, but fail to explain why most O₃ in the region is transported from the outside regions, as suggested by O₃ source apportionment. It indicates that the O₃ concentration budget cannot completely illustrate the effects of transport and photochemistry on regional O₃ pollution. By calculating, analysing and comparing the O₃ concentration and mass budget, this study aims to comprehensively understand the role of transport and photochemistry in regional O₃ pollution.

The contents of this paper includes:

- 1) Development of the method to quantify the two O₃ budgets (Sect. 2.1-2.3);
- 2) Analysis and comparison of the results from the two O₃ budgets (methodology described in Sect. 2.5, results discussed in Sect. 3);
- 3) Assessment of the role of transport and photochemistry in determining the regional origins of O₃ (methodology described in Sect. 2.6, result presented in Sect. 4).

Results show that photochemistry dominates the changes of O₃ concentrations, or plays a major role in the O₃ concentration budget. Although transport only leads to limited changes of O₃ concentrations, its large contributions in the O₃ mass budget ensure that it determines the characteristics of O₃ pollution, e.g., the regional origins of O₃ in this study. Based on the conclusions, we suggest the insights from both concentration and mass budgets are necessary to comprehensively understand the role of transport and

chemistry in regional O₃ pollution. Suggestions based on the two O₃ budgets are also provided for policy-makers when making strategies to alleviate O₃ pollution.

Our responses to specific comments and corresponding revisions are as follows (in blue and red, respectively). Note that line numbers are these in the revised manuscript with author's changes.

Specific comments:

1) Abstract: difficult to get the important of the problem from the four first lines.

Response:

We revised the first four lines in the Abstract as (in lines 20-24):

Understanding the role of transport and photochemistry is essential to mitigate tropospheric ozone (O₃) pollution within a region. In previous studies, the O₃ concentration budget has been widely used to determine the contributions of two processes to the variations of O₃ concentrations. These studies often conclude that local photochemistry is the main cause of regional O₃ pollution; however, they fail to explain why O₃ in a targeted region is primarily derived from O₃ and/or its precursors transported from the outside regions as reported by many studies of O₃ source apportionment.

2) The abstract does not help to understand the objective and the methodology approach. Ozone budget calculation and O₃ source apportionment studies seem two different types of approaches difficult to compare, so it is normal they provide different results.

Response:

We agree that different methods can give different results, but it is also important to know why they are different. For this study, the O₃ concentration budget fails to explain why most O₃ is transported from the outside regions, suggesting that this method cannot completely illustrate the effects of transport and photochemistry on regional O₃ pollution. By calculating, analysing and comparing the O₃ concentration and mass budgets, this study not only more comprehensively reveals the role of transport and photochemistry in regional O₃ pollution, but also clarifies the connections between O₃-related processes and the characteristics of O₃, i.e. the regional origins of O₃ in this study.

To make a clearer introduction, we revised the objective and the methodology in the abstract, shown in lines 27-32:

Here, we present a method to calculate the hourly contributions of O₃-related processes to the variations of not only the mean O₃ concentration, but also the total O₃ mass (the corresponding budgets are noted as the O₃ concentration and mass budget, respectively) within the atmospheric boundary layer (ABL) of the concerned region. Based on the modelling results of WRF-CMAQ, the two O₃ budgets were applied to comprehensively understand the effects of transport and photochemistry on the O₃ pollution over the Pearl River Delta (PRD) region in China.

3) Line 29: you mention two budgets, but you have not introduced them in the abstract. Is that related with the two types of studies?

Response:

Two budgets are introduced in lines 27-31:

Here, we present a method to calculate the hourly contributions of O₃-related processes to the variations of not only the mean O₃ concentration, but also the total O₃ mass (the corresponding budgets are noted as the O₃ concentration and mass budget, respectively) within the atmospheric boundary layer (ABL) of the concerned region.

The O₃ mass budget is used to explain the results of O₃ source apportionment. According to the discussions in Sect. 4 of this paper, transport and photochemistry determine the regional origins of O₃ by influencing their contributions in the O₃ mass budget as well as the regional origins of O₃ mass attributed to these O₃-related processes.

4) Line 74: the subject of that sentence “O₃ source” does not make sense. Could you elaborate more the idea in that sentence.

Response:

We agree that “O₃ source” might be a confusing item for the readers. Here, “O₃ source” was used to indicate the regional origins of O₃, or how much the concerned regions contribute to O₃ pollution. We revised the sentence into (in lines 107-111):

O₃ source apportionment is performed to identify the regional and/or sectoral origins of O₃, of which the results are also used to support air pollution control (Clappier et al., 2017; Thunis et al., 2019). Here, we only discuss the regional origins of O₃, because the contributions of sources outside the region (or emissions within the region, defined as local emissions hereafter) provide information on the influence of transport (or photochemistry) on O₃ pollution.

We also revised other “O₃ sources” in the manuscript into “the results of O₃ source apportionment”, “regional origins of O₃” or alike items.

5) Line 88: “O₃ source studies”. Use the same set of words to mention these studies. I guess in this case you want to say “O₃ source apportionment studies”. The same comment in lines 90-91, “source apportionment studies” and “O₃ budget studies”.

Response:

We accept your suggestion. However, the two sentences mentioned in the comment were deleted in the revised version. For the similar expressions afterwards, we revised them into “O₃ source apportionment studies”, “O₃ concentration budget studies” or alike items.

6) Line 93: “CTM are capable of reproducing O₃ processes”. In this sentence, you are attributing too much credibility to CTM, but models are not perfect and not always reproduce all the processes. I would be more realistic with what CTM can do, so I would suggest to rewrite this sentence.

Response:

Thanks for your suggestion. The expression here is inaccurate. In the revised manuscript, this sentence was deleted.

7) Lines 93-103 is specifically to CMAQ, it does not apply to any Eulerian CTMS (i.e. not CTM has a PA module).

Response:

Accepted. In the revised manuscript, we pointed out that the method is applied to budget calculations based on WRF-CMAQ results, as shown in lines 173-174:

WRF-CMAQ employs the Process Analysis (PA) module to assess the contributions of O₃-related processes to the variations of O₃ concentrations within each grid cell.

8) Line 121: “Horizontal transport through the borders of the PRD in four directions”. Is that correct? I guess you have two horizontal directions (x and y).

Response:

We did not state it clearly. The borders of the PRD were classified as the north, south, west, east border. Horizontal transport through four types of the PRD borders were separately quantified in O₃ budgets. Thus, we added more explanations about the classification of border (grid) in lines 199-201:

The PRD grids with one or several interfaces with the outer regions are defined as the border grids, and they can be further classified as the grids in the north, south, west and east borders based on their locations.

and also revised the expressions about the horizontal transport processes in lines 204-205:

The transport processes include horizontal transport through the four types of borders and vertical exchange through the ABL top.

9) Section 2.5 Model setup and validation. Even the model setup is described in the Qu et al. (2021) some basic details should be provided in the text, for example CMAQ and WRF version. Furthermore, the section is named “validation”. You mainly referenced Qu et al. (2021) but readers would appreciate a paragraph describing “why” we can trust on your modeling system’s results. The evaluation of ABL height with IAGOS measurements is very interesting. Could you elaborate more on the problems with CMAQ during the night?

Response:

We agreed that it is necessary to provide more details on model setup, thus relative contents were added in lines 334-352:

The WRF (version 3.2) and CMAQ (version 5.0.2) models were used to simulate the meteorological and pollutant fields, respectively. Two domains with the resolution of 36 and 12 km (denoted as d01 and d02 hereafter) were set up for the one-way nested simulations, and results in the finer d02 were used in the calculations of O₃ budgets. To represent the contributions of global background to O₃, the initial and

boundary conditions for the coarse d01 domain were provided from the global model, the Model for Ozone and Related Chemical Tracers, version 4 (MOZART-4). The PRD inventory provided by the Guangdong Environmental Monitoring Centre, the Multi-resolution Emission Inventory for China (MEIC) inventory for the mainland China (He, 2012), the MIX inventory for the Asian regions outside of mainland China (Li et al., 2017) and biogenic emissions simulated by the Model of Emissions of Gases and Aerosols from Nature (MEGAN; version 2.10) model were used in the simulations. SAPRC07 (Carter, 2010) and AERO6 were applied as the gas-phase chemistry mechanism and the aerosol scheme, respectively. The simulations of O₃ pollution in the PRD were performed for October 2015 (October 11–November 10, 2015) and July 2016 (July 1–31, 2016), which were selected as the representative months in autumn and summer, respectively. Here, O₃ polluted days are defined when the maximum hourly O₃ concentrations of the day exceed 200 µg/m³, or the maximum 8-hour average O₃ concentrations of the day exceed 160 µg/m³ (both are the Grade-II O₃ thresholds in the Chinese National Ambient Air Quality Standard) in any municipality of the PRD. According to this definition, there were 16 and 12 O₃ polluted days in the two months, respectively (more information is given in Table S1). The mean O₃ budgets during these days were calculated and discussed in the present study.

As for the validation, we agreed that the relative discussions were limited in this part. Thus, we gave more information on:

1) the validation of meteorological parameters, O₃, NO₂ concentrations and the mixing ratios of hydrocarbons by Qu et al. (2021);

2) the validation of atmospheric boundary layer height, wind speed, direction and ozone mixing ratio at different heights described in detail in Text S3

in a new paragraph, as shown in lines 354-369:

We evaluated the performance of WRF-CMAQ modelling based on multiple observational datasets. The modelling results of meteorological parameters (including temperature, relative humidity and wind speed), O₃, NO₂ concentrations and the mixing ratios of hydrocarbons were validated with corresponding observations in the PRD by Qu et al. (2021a). The performance of the model to simulate the above variables was overall satisfying with low biases and high correlations (for details, see Qu et al., 2021a). In this study, we further compared the modelled ABL height, the vertical profiles of wind speed, direction and O₃ mixing ratio in Hong Kong (located in the south PRD) with the corresponding observations from the IAGOS (In-service Aircraft for a Global Observing System; Petzold et al., 2015) dataset. The modelled ABL heights showed similar hourly variations during the day as the observational results ($R = 0.76$), with mean bias of -1.1 m (Fig. S2). The mean biases of mean wind speeds are within the range of ± 1 m/s in all height ranges (0-1 km, 1-2 km, 2-5 km), and the results of IAGOS and WRF model indicate similar variations of prevailing wind directions in different seasons and height ranges (Fig. S3). Moreover, modelled O₃ mixing ratios in Oct. 2015 are overestimated by 6% and 26% in the height range of 0-1 km and 1-2 km, respectively, and sufficiently illustrate the development, maintenance and dissipation of O₃ pollution during the month (Fig. S4). More detailed evaluations on the model performance of these parameters are presented in Text S3 of the Supplement. Overall, the model performance is acceptable, indicating that the model can provide reasonable data for the calculations of O₃ budgets.

In this study, we evaluated the modelling performance of atmospheric boundary layer (ABL) height based on the IAGOS potential temperature profiles during daytime, but not at night. One reason is that in Oct.

2015, night-time records are less ($30/105 = 28.6\%$) due to reduced flights at night. Besides, by using potential temperature profiles to determine night-time stable ABL height, large errors may occur (Dai et al., 2014). In order to have more precise O₃ budgets, more concerns on night-time ABL height are surely needed in further observations and model validation.

10) Line 220: “acceptable” from which point of view?

Response:

In this part, we evaluated the modelling performance of atmospheric boundary layer height, wind speeds, directions and O₃ mixing ratios at different heights. The results are summarized as follows:

- The modelled ABL heights showed similar hourly variations during the day as the observational results ($R = 0.76$), with mean bias of -1.1 m.
- The mean biases of mean wind speeds are within the range of ± 1 m/s in all height ranges (0-1 km, 1-2 km, 2-5 km), and the results of IAGOS and WRF model indicate similar variations of prevailing wind directions in different seasons and height ranges.
- Modelled O₃ mixing ratios in Oct. 2015 are overestimated by 6% and 26% in the height range of 0-1 km and 1-2 km, respectively, and sufficiently illustrate the development, maintenance and dissipation of O₃ pollution during the month.

High correlations and low biases of these parameters ensures that the modelling results can be used for further analyses, thus they are “acceptable”.

According the comment No. 9, relative results are described in the revised manuscript, in lines 354-369.

11) Line 221: “reasonable” from which point of view?

Response:

This question is similar as the last one. The good performance of key parameters indicates that the modelling results are close to these in reality, thus they are “reasonable” for further usage in O₃ budget calculations.

12) Line 236-237: Is that sentence well written?

Response:

We revised the sentence into (in lines 387-391):

The question to be addressed is how O₃-related processes determine the regional origins of O₃. By combining the O₃ mass budget calculations with the BFM source apportionment method, we identified the regional origins of O₃ mass changes due to transport and photochemistry (gas-phase chemistry).

13) Source apportionment method: Could you comment on the brute force disadvantages for O₃ source apportionment calculation? Could CMAQ-ISAM source apportionment method improve your results?

Response:

For this part of the study, the goal is to identify the regional origins in the O₃ mass changes attributed to transport and gas-phase chemistry (photochemistry). Besides the base scenario, three sensitivity scenarios need to be simulated in the Brute Force Method (BFM), which means increased simulation cost. But the regional source contributions in the O₃ mass changes attributed by non-transport processes, including gas-phase chemistry (photochemistry), can be identified. As a tagging method, the ISAM module in CMAQ can be used to identify the regional origins in the O₃ mass changes attributed to transport by using O₃ concentrations contributed by various regions in calculations. The simulation costs can be reduced, since it is not needed to simulate three sensitivity scenarios. However, as far as we acknowledge, the results for gas-phase chemistry (photochemistry) cannot be provided by the ISAM.

14) Conclusions: “This study concluded that transport and gas-phase chemistry play the main role in the O₃ concentration and mass budgets”. Is it not new, right? Could you elaborate more this sentence as the main conclusion of this work.

Response:

Main conclusions of this study are given in the first paragraph of Sect. 5. This paragraph aims to discuss the application of O₃ budgets in the practice of O₃ pollution control. As the first sentence, this sentence fails to start the afterward discussions, thus was revised as (in lines 726-728):

The present study concluded that transport and gas-phase chemistry play the main role in the O₃ mass and concentration budgets, respectively. As a consequence of our assessment, what should policy-makers do to effectively alleviate regional O₃ pollution?

15) Conclusions: Could you elaborate more in the biases in your modelling results? For example, discussing the uncertainties in your emission data, meteorological fields, chemical and meteorological boundary conditions, chemistry in the models.

Response:

Emissions, meteorological fields, chemical and meteorological boundary conditions, chemistry and many other factors in models could all influence the results of two O₃ budgets. However, this study focuses on the comparison between two O₃ budgets to provide a complete view on the role of transport and photochemistry in regional O₃ pollution. To have more precise O₃ budgets, we suggest to conduct more supporting observations and have more comparisons between observational and modelling results. Specifically, the observational and modelling contributions by various O₃-related processes in the O₃ budgets can be directly compared. Such results are important for further model development because it indicates which process contribute to high uncertainties in O₃ modelling. Relative contents are discussed in the Sect. 5, in lines 717-724.

Uncertainty remains in the calculated O₃ budgets, which is partly related to the biases in the modelling results. Therefore, supporting observations are essential for future research. Recent progress in observational techniques (Zhao et al., 2021; Zhou et al., 2021) has enabled three-dimensional measurements of meteorological parameters and O₃ concentrations with high spatiotemporal resolution and coverage. These data can be used not only for the model validation of key parameters in budget calculations, but also for the comparisons between observation- and modelling-based contributions by

various O₃-related processes in O₃ budgets (Kaser et al., 2017). The comparison of contributions by O₃-related processes is indicative of the main uncertainties in O₃ pollution modelling, and is therefore also important for further model developments.

Technical corrections:

1) Line 93: CTM not defined

Response:

This sentence containing “CTM” was deleted in the revised manuscript.

2) Line 95: PA module not defined.

Response:

Revised accordingly in line 173:

WRF-CMAQ employs the Process Analysis (PA) module to assess the contributions of O₃-related processes...

3) I would suggest used “tropospheric ozone” instead of “ambient O₃” when possible.

Response:

We agreed that to avoid confusion with ozone in stratosphere, “tropospheric ozone” is a better term to be used. It was revised accordingly in line 47-49:

Since first recognized as a key contributor to the Los Angeles smog, tropospheric ozone (O₃) pollution has received considerable attentions in many highly populated areas in the world...

Afterwards, “O₃” is directly used for relative discussions.

Additional statement:

Due to their strong professionalism in the areas of atmospheric pollution and modelling as well as high involvement in revising this paper, we are honoured to add Maria Kanakidou and Guy Brasseur as co-authors of this paper.

References

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Response to Reviewer II

General Comments:

Qu et al. present an analysis of the O₃ budget in the ABL in two different ways: a concentration budget and a mass budget. They apply the budget calculations to the O₃ budget over the Pearl River Delta based on simulations with WRF-CMAQ. The 2 different ways of calculating the O₃ budget lead to opposing views on the main contributions to the O₃ budget: while photochemistry dominates in the concentration budget, (vertical) transport dominates the mass budget. A tool is developed to calculate the budget contributions. A control simulation is performed, and in addition 3 brute force emission reduction scenarios are carried out. Budget calculation following the 2 methods are performed and the differences discussed.

Unfortunately, the way the paper is written makes it hard to judge its scientific merits, and I cannot recommend acceptance in its current form.

Response:

We appreciate the valuable comments and suggestions. We've tried to adjust the structure of the paper and make a lot of revisions to improve its readability.

Our responses to specific comments and corresponding revisions are as follows (in blue and red, respectively). Note that line numbers are these in the revised manuscript with author's changes.

Major comments:

1) This is a dense paper without much guidance for the reader as to where you are going, which makes it hard to follow, and hard to judge the scientific merits of the work you describe. I had to reread it 3 times and still I am getting lost in the details. Please rewrite it in a more structured way, and indicate the purpose of each section in its first sentence. For instance, in section 2.6 a number of scenario runs seems to appear out of the blue. Where are the results of these runs used/discussed?

Response:

Thanks for the suggestions. We have revised the manuscript and made it more structured, clear and reader-friendly. Pointing out the purpose of each section is surely a good way to provide readers more clues in reading — we have applied this suggestion in the revisions.

The basic logic of this paper is as follows. The objective is to comprehensively illustrate the effects of transport and photochemistry on regional O₃ pollution from the perspectives of both O₃ concentration and mass budgets. Three tasks are included in this study:

- 1) Development of the method to quantify the two O₃ budgets (Sect. 2.1-2.3);
- 2) Analysis and comparison of the results from the two O₃ budgets (methodology described in Sect. 2.5, results discussed in Sect. 3);
- 3) Assessment of the role of transport and photochemistry in determining the regional origins of O₃ (methodology described in Sect. 2.6, result presented in Sect. 4).

In the introduction part, we re-wrote the relevant paragraphs to overview the structure of this manuscript, as shown in lines 157-192:

In the ABL of the concerned region, the mean O₃ concentration and total O₃ mass are both conserved, which means that their variations are equal to the net contributions by various O₃-related processes including transport and photochemistry. These relationships can be represented by the O₃ concentration budget and mass budget, respectively. Unlike the aforementioned O₃ concentration budget in Eq. (1), the hourly O₃ mass budget, written as

$$\frac{\partial m_{O_3}}{\partial t} = -(\bar{u}s_x\langle c_{O_3} \rangle + \bar{v}s_y\langle c_{O_3} \rangle) - \overline{c_{O_3}'w'}s_z + S(O_3)V \quad (2)$$

is seldom reported (m_{O_3} is the total O₃ mass within the ABL of the region; s_x , s_y , s_z are the areas of the interfaces in the x-, y- and z-direction, respectively; V is the volume of the ABL column). Due to the varied effects of transport on O₃ concentration and mass, the O₃ mass budget differs from the O₃ concentration budget but is more suitable to explore the influence of transport and photochemistry on the results of O₃ source apportionment (more detailed explanations are given in Sect. 2.4). In order to comprehensively understand the role of transport and photochemistry in regional O₃ pollution, in the present study, we developed a method to calculate both the O₃ concentration and mass budget based on the simulation results from the Weather Research and Forecasting (WRF) and Community Multiscale Air Quality (CMAQ) models, and also analysed, compared the results of the two regional-level O₃ budgets. The Pearl River Delta (PRD) region, a city cluster located on the southeast coast of China and exposed to severe O₃ pollution in summer and autumn (Gao et al., 2018), was selected as the targeted region. The tasks for this study can be summarized as follows:

1) Development of the method to quantify the two O₃ budgets

WRF-CMAQ employs the Process Analysis (PA) module to assess the contributions of O₃-related processes to the variations of O₃ concentrations within each grid cell. However, to obtain the regional-level O₃ concentration and mass budgets, the results of PA module are not sufficient. One reason is that the contribution of vertical exchange through the ABL top is not specifically quantified in commonly used ABL parameterizations, thus requires additional calculations (Kaser et al., 2017). Additionally, calculations based on the PA results are needed to identify the contributions of other O₃-related processes to ABL-mean O₃ concentration as well as the results of the O₃ mass budget. To address this, we developed a method to quantify the two O₃ budgets, of which the details are given in Sect. 2.1-2.3.

2) Analysis and comparison of the results from the two O₃ budgets

Based on the simulations of O₃ pollution in the PRD with the model setup introduced in Sect. 2.5, the two O₃ budgets were calculated for further analyses and comparisons to reveal the role of transport and photochemistry in regional O₃ pollution from a more comprehensive perspective. Relative discussions are presented in Sect. 3.

3) Assessment of the role of transport and photochemistry in determining the regional origins of O₃

The Brute Force Method (BFM; Clappier et al., 2017), a widely used source apportionment method, was combined with the O₃ mass budget calculation to determine the contributions of emissions within and

outside the PRD as well as background sources to the O₃ transported into or produced by photochemistry in the region (methodology described in Sect. 2.6). The results, as discussed in Sect. 4, reveal the impacts of transport and photochemistry in determining the regional origins of O₃ in the PRD, and explain why the different views on the role of two processes in regional O₃ pollution are suggested by the O₃ concentration budget and O₃ source apportionment studies.

We also separated the original Sect. 3 (named *Results*) as two parts,

Sect. 3 Analyses and comparisons of O₃ concentration and mass budget

and

Sect. 4 Effects of transport and photochemistry on the regional origins of O₃

which separately discuss the results of aforementioned task 2 and 3.

2) What is actually lacking is an explanation of why 2 different budget methods give such different results. Is it mainly a boundary conditions problem? A change in mass does not lead to a change in concentration when the background concentration is similar over larger regions? Maybe it is discussed in L445-448?

Response:

The two O₃ budgets describe the conservations of O₃ concentration and mass in the atmospheric boundary layer (ABL) of the region. As introduced in Sect. 2.4, the different results of two O₃ budgets are mainly attributed to the different effects of transport on O₃ concentration and mass. When O₃ is transported into (or out of) the ABL of the region through the advection process (horizontal transport and vertical exchange through the ABL top due to large-scale air motion (ABLex-M)), surely total O₃ mass increases (or decreases). However, whether O₃ concentration increases or decreases also depends on the difference between O₃ concentrations in the region and transported air parcels — that is why clean (polluted) air parcels being transported into the region dilutes (aggravates) O₃ pollution. The effect of transport can be understood as to replace a part of air mass with the transported air parcel. If O₃ concentration is higher (or lower) in the transported air parcel, by replacing, mean O₃ concentration within the region will increase (or decrease). This effect also applies to the exchange through the ABL top due to the temporal changes of ABL heights (ABLex-H). For example, after sunrise, O₃ mass in the ABL of the region increases rapidly along with the development of ABL. This process can be viewed as two air parcels combining into one, and whether O₃ concentration increases or decreases also depends on the difference of O₃ concentrations in two air parcels — but O₃ mass surely increases. More detailed contents are discussed in Sect. 2.4, in lines 301-321:

The difference between the two O₃ budgets is linked to the varied effects of transport on O₃ mass and concentration. Suppose that the mean O₃ concentration in the transported air parcels is $\langle c_{O_3} \rangle_{trans}$. For horizontal transport, its contributions in the O₃ mass and concentration budgets can be separately written as:

$$F_{htrans} = \langle c_{O_3} \rangle_{trans} dV \quad (8)$$

$$d\langle c_{O_3} \rangle_{htrans} = \frac{dV}{V} (\langle c_{O_3} \rangle_{trans} - \langle c_{O_3} \rangle) \quad (9)$$

Apparently, F_{htrans} is related to the O_3 concentrations in the transported air parcels, but not to those in the studied region. It indicates how much O_3 is transported into or out of the region. Whether it is positive or negative only depends on the direction of transport — O_3 being transported into (out of) the region leads to the increase (decrease) of O_3 mass, which corresponds to a positive (negative) contribution in the O_3 mass budget. In contrast, $d\langle c_{O_3} \rangle_{htrans}$ quantifies how much horizontal transport alters regional-mean O_3 concentrations, and is linked to the difference between O_3 concentrations in the transported air parcels and the studied region (Eq. (9)). O_3 being transported into (out of) the region does not necessarily result in a higher (lower) O_3 concentration. For instance, when clean air parcels with relatively low O_3 levels are transported into the region, they dilute O_3 pollution and reduce O_3 concentration ($d\langle c_{O_3} \rangle_{htrans} < 0$). Given that ABLex-M is also an advection process, the above difference applies to this process as well. For ABLex-H, its contributions in the O_3 mass and concentration budgets are expressed as:

$$F_{ABLex-H} = \langle c_{O_3} \rangle_{trans} dV \quad (10)$$

$$d\langle c_{O_3} \rangle_{ABLex-H} = \frac{dV}{V + dV} (\langle c_{O_3} \rangle_{trans} - \langle c_{O_3} \rangle) \quad (11)$$

Similarly, ABL development and collapse lead to the increase and decrease of O_3 mass, respectively, but whether they contribute to higher or lower O_3 concentration also depends on the difference between O_3 concentration in the transported air parcels and that in the region. Based on the above discussion, these transport processes all show different effects on O_3 mass and concentration — the effect of transport on the variations of O_3 mass is only related to the characteristics of the transported air parcels, namely their volumes and O_3 concentrations within (Eqs. (8) and (10)), while how transport contributes to the variations of O_3 concentration is linked to the difference between O_3 concentrations in the transported air parcels and the region (Eqs. (9) and (11)).

It is possible that change in mass does not lead to a change in concentration when the background concentration is similar over larger regions. For example, suppose that air parcels with the volume of dV and the O_3 concentration of $\langle c_{O_3} \rangle_{trans}$ are transported into the region through the ABLex-H process. The contributions of such a process to O_3 mass and concentration (denoted as $F_{ABLex-H}$ and $d\langle c_{O_3} \rangle_{ABLex-H}$, respectively) can be expressed as (Eqs. (10-11) of the manuscript):

$$F_{ABLex-H} = \langle c_{O_3} \rangle_{trans} dV$$

$$d\langle c_{O_3} \rangle_{ABLex-H} = \frac{dV}{V + dV} (\langle c_{O_3} \rangle_{trans} - \langle c_{O_3} \rangle)$$

where V is the original volume of the ABL of the region, $\langle c_{O_3} \rangle$ is the initial mean O_3 concentration. $F_{ABLex-H}$ is surely positive, since $\langle c_{O_3} \rangle_{trans}$ and dV are both above 0. As an extreme case, if $\langle c_{O_3} \rangle_{trans} = \langle c_{O_3} \rangle$, then $d\langle c_{O_3} \rangle_{ABLex-H} = 0$, which means that the transport, or “combination”, of air parcels with the same O_3 concentration leads to increased O_3 mass and volume in the ABL of the region at the same time, but O_3 concentration does not change.

In L445-448 of the original manuscript, we discussed why the conclusions in this paper are important for further studies. Differences in the concentration and mass budgets apply to not only O_3 , but also other pollutants with moderately long atmospheric lifetimes, such as fine particulate matter and some of its components. Transport may fail to notably alter pollutant concentration, but can significantly contribute to the changes of pollutant mass. Specifically, massive pollutant being transported into the ABL in the morning nearly determines the characteristics of pollutant within the region — besides the origins of

pollutants, they also include the contributions of different reaction pathways and sensitivities to precursor emissions. But in the concentration budget, the effects of transport on these characteristics are often ignored. In order to fully understand the effects of transport, chemistry and other related processes, we suggest that the insights from both concentration and mass budgets are required for future studies.

Minor comments:

1) L50 (and throughout MS): O₃ processes --> O₃-related processes

Response:

Accepted and revised as suggested.

2) L74: pls rephrase sentence

Response:

This sentence is revised into (in lines 107-109):

O₃ source apportionment is performed to identify the regional and/or sectoral origins of O₃, of which the results are also used to support air pollution control (Clappier et al., 2017; Thunis et al., 2019).

3) L416: “High contributions of ...” Unclear sentence. Please rephrase.

Response:

This sentence is revised into (in lines 650-654):

By combining the O₃ mass budget and O₃ source apportionment, we identified the O₃ mass increase due to O₃-related processes as local (PRD) and non-local (EC-China and BCON) contributions. According to the results discussed before, high contributions of transport in the morning-hour O₃ mass increase and the dominance of non-local source contributions in this part of new O₃ ensure that non-local sources contributed to most O₃ in the PRD.

4) L461: what do you mean by ‘a longer time’?

Response:

“A longer time” is vague, thus it is revised as in lines 732-733:

However, for short-term air pollution control, this strategy is not efficient because emission reduction in upwind regions may need to start days earlier before the polluted periods.

Additional statement:

Due to their strong professionalism in the areas of atmospheric pollution and modelling as well as high involvement in revising this paper, we are honoured to add Maria Kanakidou and Guy Brasseur as co-authors of this paper.

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