

## ***Interactive comment on “Impacts of aerosols on the chemistry of atmospheric trace gases: a case study of peroxides and HO<sub>2</sub> radicals” by H. Liang et al.***

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Thanks for your constructive and thoughtful comments. We have revised our manuscript, according to your comments. Below is our response, as shown in answer (A), to your comments.

General: This is a sound study on peroxides, mainly studied in field measurements in Beijing, China which in a modelling part also deals with the impact of aerosol particles on gas phase HO<sub>2</sub> and hence H<sub>2</sub>O<sub>2</sub>. The field measurement part contains a number of interesting results. The modeling part much resembles recent work by Mao et al.

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Overall, the paper can be published in ACP subject to a few revisions.

A: Thank you for your constructive comments.

Page 16557, line 7: The model mechanism which has been used should be fully documented. It is not sufficient to state "it has been chiefly drawn"....The full documentation can be given in a supplement.

A: Yes, we have given the full documentation as a table in the Supplement (Table S1).

P 16562, l 24: Schuchmann and von Sonntag

A: Yes, we have corrected it.

P 16572, l 15ff: Please give evidence that really diffusion is a very important difference between aerosol and haze periods. That might also be due to different aqueous phase chemical conversion as concentration change.

A: Yes. We think that the aqueous phase diffusion of HO<sub>2</sub> is really important for the differences between non-haze and haze periods. According to Jacob (2000), the corresponding characteristic time of aqueous phase diffusion and chemical reactions in aerosols could be given as follows:

$$\tau_{diff} = \frac{r^2}{\pi^2 D_l}$$
$$\tau_{chem} = \frac{1}{k_{chem}}$$

where  $r$  is the radius of the particle (m),  $D_l$  is the aqueous phase diffusion coefficient ( $\text{m}^2 \text{s}^{-1}$ ), and  $k_{chem}$  is the first-order reaction rate constant. For HO<sub>2</sub>, both  $\tau_{diff}$  and  $\tau_{chem}$  are on the order of  $10^{-6}$ -  $10^{-5}$  s, indicating that both processes could affect the uptake coefficient of HO<sub>2</sub>. Larger particle radius in the haze period would lead to a slower aqueous phase diffusion and hence a smaller  $\gamma_{HO_2}$ . We agree that in addition to diffusion,  $\gamma_{HO_2}$  is affected by the different chemical conversions due to the HO<sub>2</sub> concentration change. We have rewritten the relevant statement in the revised manuscript as follows: "We estimated the characteristic times of aqueous phase diffusion ( $\tau_{diff}$ )

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and chemical reactions ( $\tau_{chem}$ ) for HO<sub>2</sub> according to the equations given by Jacob (2000), and found that both  $\tau_{diff}$  and  $\tau_{chem}$  were on the same order. This indicates that  $\gamma_{HO_2}$  would be determined by the combination of both processes.”

#### References

Jacob, D. J.: Heterogeneous chemistry and tropospheric ozone, *Atmos. Environ.*, 34, 2131-2159, 10.1016/S1352-2310(99)00462-8, 2000.

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