## Dear Editor,

Thank you for your recommendation to respond more clearly on the reviewer's #3 comment about used in our work rate coefficient of the reaction of  $SO_2$  with OH. Following this recommendation, we extended the Table 1 with the results showing the contribution of the OH+SO<sub>2</sub> reaction if the lower rate coefficient from Medeiros et al. (2018) is used.

The Table 1 was modified as follows (changes are shown in red):

**Table 1.** Comparison of observed  $[H_2SO_4]$  with calculations assuming  $H_2SO_4$  formation via oxidation of  $SO_2$  by OH and SCIs. [OH] and  $[H_2SO_4]$  are observed concentrations,  $[H_2SO_4]^{OH}$  and  $[H_2SO_4]^{SCI}$  are calculated  $H_2SO_4$  produced by oxidation of SO<sub>2</sub> by OH and SCI, respectively. Values in square brackets correspond to  $[H_2SO_4]^{OH}$  concentrations calculated with a rate coefficient for the reaction of OH with SO<sub>2</sub> from Medeiros et al. (2018) (see discussion in Sect. 4.2).

	Daytime: 7:00 – 20:00		Night-time: 20:00 – 7:00	
	Median (inter-quartile range)	$Mean \pm 1\sigma$	Median (inter-quartile	Mean $\pm 1\sigma$
			range)	
$[OH], 10^5 \text{ cm}^{-3}$	31 (18; 42)	$31 \pm 17$	1.1 (-0.7; 3.0)	$1.7\pm4.0$
$[H_2SO_4], 10^5 \text{ cm}^{-3}$	47 (28; 86)	$63 \pm 49$	4.2 (3.1; 6.4)	$5.8\pm4.8$
$[H_2SO_4]^{OH} = a + b \times [H_2SO_4]$	$a=(-2.0\pm0.5)\times10^5$ ; $b=0.85\pm0.02$		$a=(-3.1\pm0.3)\times10^5$ ; b=0.97±0.1	
	$[a=(-1.1\pm0.3)\times10^5; b=0.52\pm0.01]$		$[a=(-1.7\pm0.2)\times10^5; b=0.60\pm0.05]$	
$[H_2SO_4]^{OH/}$ $[H_2SO_4]$ , %	95 (79; 129)	$86 \pm 4$	39 (-8; 84)	$9\pm15$
	[58 (48; 78)]	$[52 \pm 2]$	[23 (-4; 52)]	[5 ± 9]
1-[H <sub>2</sub> SO <sub>4</sub> ] <sup>OH</sup> / [H <sub>2</sub> SO <sub>4</sub> ], %	5 (-29; 21)	$14 \pm 4$	61 (16; 108)	$91\pm15$
	[42 (22; 52)]	$[48 \pm 2]$	[77 (48; 104)]	$[95 \pm 9]$
$[H_2SO_4]^{SCI}/[H_2SO_4], \%$	10 (7; 16)	$12\pm 6$	30 (22; 48)	$38\pm24$
[H <sub>2</sub> SO <sub>4</sub> ]-[H <sub>2</sub> SO <sub>4</sub> ] <sup>OH</sup> , 10 <sup>5</sup> cm <sup>-3</sup>	1.2 (-12.5; 8.1)	$4.6\pm3.2$	3.0 (0.8; 5.2)	$3.1 \pm 0.4$
	[15.2 (8.3; 36.5)]	$[11.2 \pm 3.4]$	[3.2 (1.8; 5.2)]	$[3.3 \pm 0.3]$
$[H_2SO_4]^{SCI}$ , 10 <sup>5</sup> cm <sup>-3</sup>	6.0 (3.7; 8.6)	$6.4\pm3.7$	1.4 (1.1; 2.4)	$1.8 \pm 1.2$

Also, we have added some comments to the Sect. 4.2, where this issue is discussed:

"In a recent study of Blitz et al. (2017a, 2017b) a significantly lower rate constant of  $5.8 \times 10^{-13}$  cm<sup>3</sup> molecule<sup>-1</sup> s<sup>-1</sup> was derived from experiments with vibrationally excited OH (v=1,2,3)+SO<sub>2</sub> and using the master equation analysis of the pressure and temperature dependence of their own and some others experimental OH+SO<sub>2</sub> reaction rate constants. An even lower rate constant of  $4.8 \times 10^{-13}$  cm<sup>3</sup> molecule<sup>-1</sup> s<sup>-1</sup> has been derived by Medeiros et al. (2018) applying more detailed master equation analysis of experimental data from Blitz et al. (2017a, 2017b) and some other data. These recent results have not been confirmed by other studies. Also, they have been discussed but not recommended by the latest JPL evaluation (Burkholder et al., 2019).

Using the lower rate constant from Medeiros et al. (2018) in our study would result in about 2 times reduced  $H_2SO_4$ production by oxidation of  $SO_2$  by OH and would invoke either significantly larger contribution from an additional  $H_2SO_4$  source or a lower  $H_2SO_4$  uptake coefficient, of about 0.5 instead of unity. As shown in Table 1, the reaction of OH with  $SO_2$  would explain only about 50% and 5% of the observed  $H_2SO_4$  production during the day and during the night, respectively."

Sincerely, Alexandre Kukui