

## ***Interactive comment on “The role of volatile organic compounds in the polluted urban atmosphere of Bristol, UK” by A. C. Rivett et al.***

**A. C. Rivett et al.**

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The authors would like to thank the referee for his careful comments. The points raised are answered below. Referee's comments are shown in bold.

### **Technical comments      1-5, 8-10, 14, 18-21**

Altered in text as suggested.

**6)** Please see the response to a similar question in the other Referees Report (acpd-3-S369)

### **7) Page 774, an example of exactly how the formula (1) was used**

A calculation of daytime [OH] using Propene as an example:

$$[OH] = \frac{\ln((X_t/B_t)/(X_i/B_i))}{t(K_B - K_X)}$$

$$X_i/B_i = 0.769533359$$

$$X_t/B_t = 0.363993344$$

$$K_X = 2.70 \text{ E-11 cm}^3 \text{ molecules}^{-1} \text{ s}^{-1}$$

$$K_B = 1.32 \text{ E-12 cm}^3 \text{ molecules}^{-1} \text{ s}^{-1}$$

$$t = \text{Time between morning rush hour and midday periods} = 16\,200 \text{ s}$$

The ratio of propene to benzene ( $X_i/B_i$  and  $X_t/B_t$  during the morning rush hour period (7:00-10:00) and the midday period (10:00-16:00) respectively) is the slope from a graph of propene concentration plotted against benzene concentration during those periods, using data from over the whole campaign.  $K_X$  and  $K_B$  are the rate constants for the reaction of benzene and hydrocarbon  $X$  with OH.

$$[OH] = \frac{\ln(0.364/0.770)}{16\,200(1.32 \times 10^{-12} - 2.70 \times 10^{-11})}$$

$$[OH] = \frac{\ln(0.473)}{16\,200(-2.568 \times 10^{-11})}$$

$$[OH] = \frac{-0.749}{-4.16 \times 10^{-7}}$$

Calculated Daytime OH Concentration =  $1.80 \times 10^6 \text{ molecules cm}^{-3}$

**"...which rate constant for benzene that was used (with reference), which results came out of the formula and what was the conclusion (Table 1?)."**

Rate constants are given in Appendix 2 and the results in Table 1. The conclusions we drew were... "The average daytime OH concentration calculated over the course of the campaign was  $2 \times 10^6 \text{ molecules cm}^{-3}$ . This compares well with what would be expected, as the lifetime of OH is less than 1 second and consequently

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noontime concentrations generally only reach 0.04-0.4 ppt ( $10^6$ - $10^7$  molecules  $\text{cm}^{-3}$ ) (Wayne, 2000)."

**11) Page 776, formula (6): How do you calculate/know the dispersion factor?**

The dispersion factor is calculated from the hydrocarbon decays using Equation 12.

**12)** The complicated series of equations in Appendix 3 are derived from Equation 6 and these are then used to calculate radical concentrations and the dispersion factor.

**13) Page 776, line 12: What is the uncertainty of the value  $3.8 \times 10$  to the 5 molec  $\text{cm}^{-3}$ ?**

The uncertainty in the estimated night-time OH value is the range of concentrations calculated, which are given.

**15) Page 778, Why are isoprene and the monoterpenes not included in the calculations?**

These species were unfortunately not measured during the campaign. Have added an explanation of this.

**16) + 17)** The photochemical ozone creation potentials for each of the hydrocarbons are calculated in the following way:

$$\text{POCP} = (\text{Hydrocarbon Concentration} / \text{Propane Concentration}) \times \varepsilon^{\text{POCP}}$$

$$\text{POCP} = (\text{Hydrocarbon Concentration} / \text{Propane Concentration}) \times \gamma_R$$

The  $\varepsilon^{\text{POCP}}$  and  $\gamma_R$  indices used in the calculations are given in Appendix 4 and were taken from Jenkin, 1998 and not calculated by the authors.

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Interactive comment on Atmos. Chem. Phys. Discuss., 3, 769, 2003.

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