

Peroxy radicals during BERLIOZ at Pabstthum: Measurements, radical budgets and ozone production

D. Mihelcic,¹ F. Holland,¹ A. Hofzumahaus,¹ L. Hoppe,¹ S. Konrad,¹ P. M $\ddot{u}$ sgen,¹ H.-W. P $\ddot{a}$ tz,¹ H.-J. Sch $\ddot{a}$ fer,¹ T. Schmitz,¹ A. Volz-Thomas,¹ K. B $\ddot{a}$ chmann,² S. Schlomski,² U. Platt,³ A. Geyer,³ B. Alicke,³ and G. K. Moortgat⁴

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[1] The Photochemistry Experiment during BERLIOZ (PHOEBE) was conducted in July and August 1998 at a rural site located near the small village of Pabstthum, about 50 km northwest of downtown Berlin. In this paper, spectroscopic measurements of hydroxyl (OH) and peroxy radicals (HO₂ and RO₂) are discussed for two intensive days (20 and 21 July) of the campaign. On both days peak values of the radical concentrations were similar, reaching $6\text{--}8 \times 10^6 \text{ cm}^{-3}$ for OH and 20–30 ppt for RO₂ and HO₂. Fairly high OH concentrations were observed during the morning hours in the presence of high-NO_x mixing ratios (>20ppb). The “master chemical mechanism” (MCM) was used to calculate OH, HO₂, and RO₂ concentrations from the simultaneously measured data comprising a comprehensive set of speciated hydrocarbons and carbonyl compounds, O₃, CO, NO, NO₂, HONO, PAN, J(NO₂), J(O¹D), and meteorological parameters. The calculated OH concentrations are in excellent agreement with the measurements during the morning hours at high-NO_x (>10 ppb). However, at low NO_x conditions the model overestimates OH by a factor 1.6. The modeled concentrations of HO₂ and RO₂ are in reasonable agreement with the measurements on 20 July. On the next day, when isoprene from nearby sources was the dominant VOC, the model overpredicted HO₂ and RO₂ in addition to OH. Radical budgets solely calculated from measured data show that a missing sink for OH must be responsible for the overestimation by MCM. Missing VOC reactivity is unlikely, unless these VOC would not lead to RO₂ production upon reaction with OH. The measured RO₂/HO₂ ratio of about one is well reproduced by the MCM, whereas a simple model without recycling of RO₂ from decomposition and isomerisation of alkoxy radicals underpredicts the measured ratio by about a factor of two. This finding highlights the importance of RO₂ recycling in the chemical mechanism. The ozone production rate P(O₃), calculated from the peroxy radical concentrations and NO, had a maximum of 8 ppb/hr at 0.5 ppb NO, which is in good agreement with results from previous campaigns at Tenerife and Schauinsland. **INDEX TERMS:** 0365 Atmospheric Composition and Structure: Troposphere—composition and chemistry; 0345 Atmospheric Composition and Structure: Pollution—urban and regional (0305); 0322 Atmospheric Composition and Structure: Constituent sources and sinks; **KEYWORDS:** hydroxyl radical, peroxy radicals, radical budgets, ozone production, photochemistry

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1. Introduction

[2] The hydroxyl radical (OH) is the major tropospheric oxidant and is responsible for the chemical degradation of

many atmospheric trace gases and pollutants [Levy, 1971; Logan et al., 1981; Ehhalt et al., 1991]. Peroxy radicals (HO₂ and RO₂, where R represents an organic group) are produced by reactions of OH with CO and volatile organic compounds (VOC) and play a central role in the photochemical formation of ozone in the troposphere [Chameides and Walker, 1973; Crutzen, 1979; Trainer et al., 1987].

[3] In recent years, measurements of OH, HO₂, and the sum of HO₂ + RO₂ have become available in a number of field studies, which were designed to test our understanding of the fast photochemistry in the planetary boundary layer. The experiments were performed at different sites in clean Antarctic [Jefferson et al., 1998; Chen et al., 2001] and

¹Institut für Chemie und Dynamik der Geosphäre, Forschungszentrum Jülich, Jülich, Germany.

²Technische Hochschule Darmstadt, Darmstadt, Germany.

³Ruprecht-Karls-Universität Heidelberg, Institut für Umweltphysik, Heidelberg, Germany.

⁴Max-Planck-Institut für Chemie, Division of Atmospheric Chemistry, Mainz, Germany.

marine environments [e.g., *Hauglustaine et al.*, 1999; *Carslaw et al.*, 1999; *Kanaya et al.*, 2000; *Brauers et al.*, 2001], in clean continental regions [*Poppe et al.*, 1994; *McKeen et al.*, 1997; *Ehhalt*, 1999;], and in forested areas [*Carslaw et al.*, 1999; *Tan et al.*, 2001]. One study has been reported which was performed in polluted urban air [*George et al.*, 1999].

[4] In only few of the ground-based studies both OH and HO₂ were measured. In the Tropospheric OH Photochemistry Experiment (TOPHE) carried out in the Rocky Mountains of Colorado in 1993, OH concentrations were measured by long-path absorption and by ion-assisted mass spectrometry [*Mount and Williams*, 1997; *Tanner et al.*, 1997]. HO₂ was measured by conversion into OH, which was subsequently detected by laser-induced fluorescence [*Stevens et al.*, 1997], and the sum of HO₂ and RO₂ was measured by a chemical amplifier [*Cantrell et al.*, 1997]. In this study the chemical model overpredicted the OH concentrations by 50% on average [*McKeen et al.*, 1997], and the modeled HO₂ concentrations were higher than the measurements by up to a factor of 3 [*Cantrell et al.*, 1997]. The measured ratio RO₂/HO₂ was 3–4 times greater than expected under clean conditions, and over a factor of 10 higher than predictions under polluted conditions [*Stevens et al.*, 1997].

[5] In the Los Angeles Free Radical Experiment (LAFRE) OH and HO₂ concentrations were measured in polluted urban air by the FAGE technique in 1993 [*George et al.*, 1999]. The observations were compared with model calculations, which used a surrogate species chemical mechanism. Early and late in the day, the model yielded good agreement with the measured OH and HO₂ concentrations. However, during midday the model results were too high by a factor of 1.5–2. Using the measured HO₂ concentrations as model input resulted in an improved agreement between modeled and measured OH.

[6] The Eastern Atlantic Summer Experiment (EASE 96) was performed in the marine environment at the Mace Head Atmospheric Observatory in 1996. OH and HO₂ were measured by the FAGE (fluorescence assay by gas expansion) technique and the sum of RO₂ + HO₂ was recorded by a chemical amplifier [*Carslaw et al.*, 1999]. The measurements were compared with the results of a model [*Carslaw et al.*, 1999] based on the detailed Master Chemical Mechanism (MCM) by *Jenkin et al.* [1997]. For OH the model predictions were too high by about 40%. For HO₂ the predictions were in good agreement with the measurements on one day, but were greater than the observations by as much as a factor of 6 on another day. For RO₂ + HO₂ a general good agreement was found between the measured and modeled concentrations over a wide range of conditions.

[7] In the Photochemistry, Emissions, and Transport (PROPHET) experiment OH and HO₂ were measured by the laser-induced fluorescence technique in northern Michigan in 1998 [*Tan et al.*, 2001]. The measurement site was located in a deciduous forest marked by relatively low NO_x levels and high isoprene emissions [*Carroll et al.*, 2001]. The measured OH and HO₂ concentrations were compared to the results of a model [*Tan et al.*, 2001], which was derived from the Regional Acid Deposition Mechanism (RADM) [*Stockwell et al.*, 1997] and modified to include

the oxidation of isoprene and other biogenic VOC. Unlike in the TOHPE, LAFRE, and EASE96 campaigns, the modeled OH concentrations were generally smaller, rather than higher, when compared to the observed OH. On average the model under prediction was a factor 2.7. For HO₂ the model results were in good agreement with the measurements. Thus modeled HO₂/OH ratios were 2.5–4 times higher than observed.

[8] In the present paper we report measurements and model calculations of OH, HO₂, and RO₂ from a local photochemistry study (PHOEBE), which was performed in the Berlin Ozone experiment (BERLIOZ) in summer 1998. BERLIOZ was a major activity in the area around the city of Berlin (Germany) and aimed to study the transport and chemical processes which control the formation of ozone and photooxidants in the city plume. At Pabstthum, which was one of several measurement sites in the Berlin area, comprehensive in situ measurements of radicals (OH, HO₂, RO₂, and NO₃) were performed, in order to study their fast chemistry in a rural environment under the influence of urban pollution [*Volz-Thomas et al.*, 2003a]. It is the first field experiment where all radicals were measured by at least one direct spectroscopic technique. Along with the radical data, a comprehensive set of measurements of NO, NO₂, NO_y, HONO, PAN, O₃, H₂O, H₂O₂, ROOH, CO, VOC, SO₂, and photolysis frequencies among them J(O¹D) (O₃ → O(¹D)) and J(NO₂) (NO₂ → NO) were obtained. The speciated VOC measurements include alkanes, alkenes, isoprene, α-pinene, ethyne, aromatics, formaldehyde, other carbonyl compounds, and carboxylic acids. In the following we focus our analysis on the daytime chemistry of OH, HO₂, and RO₂, whereas the nighttime chemistry, including reactions of NO₃, is presented in a companion paper by *Geyer et al.* [2003]. The daytime radical concentrations and their ratios are compared for two days (20 and 21 July) with model results, using an updated master chemical mechanism (MCM). The two days were characterized by high-NO_x (up to 30 ppb) in the morning and high isoprene levels (several hundred ppt) on the second day. Unlike in previous studies, speciated radical data from the measurements are used to investigate the radical budgets and the ozone production rate. The ozone formation is compared with field data obtained earlier at other sites (Tenerife, Schauinsland) in Europe.

2. Experimental Methods

[9] The PHOEBE measurement site was located in a rural area near the small village Pabstthum (52.9°N, 12.9°E) at about 50 km in the North-West of the city center of Berlin. The station was set up on the western border of a flat grassland (7.5 × 4.5 km²). A mixed forest dominated by Scots pine extended from the station in westerly and northerly direction. A description of the location, the arrangement of the instruments and details of the measurements can be found in the overview paper [*Volz-Thomas et al.*, 2003a] and in the other papers of this special section.

2.1. Peroxy Radical Measurements

[10] The peroxy radicals were measured by Matrix Isolation and Electron Spin Resonance Spectroscopy (MIESR). The method is described in detail by *Mihelcic et al.* [1985,

Table 1. Chemical Species and Photolysis Frequencies Used as Model Input Parameters or for the Radical Budget Calculations^a

Parameter	Instrument	Time Resolution	Institution	Reference
<i>Odd Nitrogen Compounds</i>				
NO	chemiluminescence (770 Alppt, EcoPhysics)	1 min	FZJ	A
NO ₂	CLD 770 Alppt + PLC 760 (EcoPhysics)	1 min	FZJ	A
HNO ₂	DOAS 1	30 min	IUP	F
PAN	GC-ECD (Meteorologieconsult)	6 min	FZJ	A
<i>Other</i>				
O ₃	UV-Absorption (TE-49)	1 min	FZJ	A
CO	IR-Absorption (TE-48)	1 min	FZJ	A
H ₂	not measured; 500 ppb			
Aerosol	Pallas Optical Counter (PCS 2000)	5 min	IUP	G
<i>Organic Compounds</i>				
CH ₄	not measured; 1.8 ppm			
NMHC: C ₅ –C ₁₀	In situ-GC (Airmotec HC1010)	20 min	FZJ	B
NMHC: C ₂ –C ₁₀	in situ-GC (HPLC/Cryogenic sampling)	85 min	FZJ	B
H ₂ CO	fluorescence (AeroLaser, AL4001)	1 min	FZJ	A
Carbonyl compounds	derivatisation, GC/ECD	1 h	TUD	E
Carboxylic acids	capillary electrophoresis-LIF	1 h	TUD	E
<i>Free Radicals</i>				
OH	LIF	90 s	FZJ	D
HO ₂	LIF	90 s	FZJ	D
HO ₂	MIESR	30 min	FZJ	C
RO ₂	MIESR	30 min	FZJ	C
<i>Meteorology and Radiation</i>				
T, rH, P, wind vector	psychrometer, wind vane, cup anemometer (Thiess)	1 min	FZJ	A
J(O ¹ D), J(NO ₂)	2π sr Filter radiometers	10 s	FZJ	D

^aReferences are as follows: A, Volz-Thomas et al. [2003a]; B, Konrad et al. [2003]; C, this paper; D, Holland et al. [2003]; E, Grossmann et al. [2003]; F, Aliche et al. [2003]; G, Aliche (personal communication, 1999). FZJ, Institut für Chemie und Dynamik der Geosphäre, Forschungszentrum Jülich; IUP, Institut für Umweltphysik, Universität Heidelberg; MPI, Max Planck Institut für Chemie, Mainz; TUD, Institut für Chemie, Technische Universität Darmstadt.

1990]. Briefly: During the sampling interval of 30 min, ambient air (total volume: 8 l STP) is expanded at a flow rate of 270 ml/min (STP) through a nozzle into a vacuum chamber. The radicals are trapped from the gas jet in a polycrystalline D₂O matrix which is formed in situ on a cold finger at a temperature of 77 K. Speciation and quantification of the radicals is achieved in the laboratory on a Bruker ESP 300E ESR-spectrometer equipped with an ER 4109 WZS wide bore cavity. The samples are kept under vacuum at 77K during storage and during the measurement. The ESR-spectra are analyzed by a nonlinear fitting procedure [Mihelcic et al., 1990]. The MIESR method allows the speciation of HO₂, CH₃C(O)O₂ and the sum of organic peroxy radicals. The latter is denoted RO₂ in the following. The MIESR also provides measurements of NO₂ and NO₃.

[11] Over the range of concentrations encountered during BERLIOZ, the precision for NO₂, NO₃, HO₂, CH₃C(O)O₂, was always better than 2 ppt. The precision for NO₃ radicals was better than 1 ppt, because of its narrower line width. The systematic uncertainty of the MIESR measurements was estimated from absolute calibrations made in the laboratory to be ±5% [Mihelcic et al., 1985, 1990; Geyer et al., 1999, 2003].

[12] The MIESR sampler was mounted on a pneumatic mast, using a ball bearing and a wind-vane to ensure that the inlet nozzle pointed into the wind in order to avoid contamination or losses of the radicals on the surface of the sampler. The inlet of the sampler was 11.5 m above the surface during sampling. Due to the limited number of samples that could be stored for later analysis, measurements by MIESR were concentrated on the intensive

observational periods which took place on 20–21 July and in the night from 4 to 5 July 1998.

2.2. Other Radical Measurements

[13] The total peroxy radical concentration (HO₂ + RO₂) was also measured at Pabstthum by a Chemical Amplifier (CA) deployed by the Max-Planck Institute for Chemistry, Mainz [Volz-Thomas et al., 2003b]. The concentrations of OH and HO₂ radicals were measured by Laser Induced Fluorescence (LIF) [Holland et al., 2003]. The LIF instrument sampled ambient air by gas expansion through a nozzle at about 10 m above ground. OH was measured directly by resonance fluorescence following excitation at 308 nm. HO₂ was first converted chemically into OH, which was subsequently detected by LIF. The time resolution of the measurements was 1.5 min and the estimated calibration error ±20% (2σ).

2.3. Other Trace Gases and Parameters

[14] A comprehensive set of trace gases, photolysis frequencies, and meteorological parameters was measured during PHOEBE [cf., Volz-Thomas et al., 2003a] (Table 1). The parameters used in this paper for model initialization and radical budget calculations are listed in Table 1 together with the appropriate references. NO and NO₂ were measured by chemiluminescence and a photolytic converter (accuracy 5% and 5% for NO and NO₂, respectively). O₃ and CO concentrations were measured using two identical standard UV-photometers (TE-49) and a slightly modified infrared filter correlation instrument (TE-48), respectively. Peroxyacetyl nitrate (PAN) was measured by gas chroma-

tography with an electron capture detector (accuracy < 10%). Nitrous acid (HONO) was measured using Differential Optical Absorption Spectroscopy (DOAS; limit of detection: 80 ppt; accuracy $\pm 6\%$, 1σ). More than 60 hydrocarbons (HC) in the range of C_2 – C_{10} were measured by using two in situ GC systems [Konrad *et al.*, 2003]. The measurements included alkanes, alkenes, isoprene, α -pinene, ethyne, and aromatic compounds. Combining the data sets from the two instruments NMHC records with a time resolution of 20 min were obtained. Carbonyl compounds (C_2 to C_{10}) were measured by a novel sampling technique, followed by analysis *via* gas chromatography. The data sets provide information on a large number of aliphatic and olefinic alkanals, including hydroxycarbonyls and dicarbonyls, benzaldehyde and pinonaldehyde [Grossmann *et al.*, 2003]. Formaldehyde was measured using a fluorescence instrument (AeroLaser) based on the Hantzsch reaction and by the DOAS instrument. Photolysis Frequencies of $O_3 \rightarrow O(^1D)$, $J(O^1D)$, and $NO_2 \rightarrow NO$, $J(NO_2)$, were measured by 2π sr filter radiometers and an actinic flux spectral radiometer with good agreement between the instruments [Holland *et al.*, 2003]. Due to data gaps caused by occasional malfunctioning of the spectral radiometer, $J(O^1D)$ and $J(NO_2)$ data are taken from the filter radiometers. The estimated errors are $\pm 14\%$ and $\pm 9\%$, respectively. Wind, temperature, pressure and relative humidity (rH) were measured by standard meteorological devices.

[15] All in situ measurements were made at approximately the same height as the MIESR, about 10–11 m above ground and within a horizontal distance of 20 m. The DOAS instrument employed a folded light path of 15 m base length [Alicke *et al.*, 2003]. It was mounted in an open crane cantilever on an 8 m high platform, next to the LIF instrument container (see Figure 3 of Volz-Thomas *et al.* [2003a] for details).

[16] We like to note that several parameters (e.g., HO_2 , NO_2 , O_3 , H_2CO , $J(NO_2)$) were measured by more than one technique. In all cases good agreement was observed between the different instruments [Volz-Thomas *et al.*, 2003a], except for formaldehyde where an unresolved discrepancy between Hantzsch and DOAS was observed with the DOAS data being systematically larger by up to a factor of two (1.3 on average) [Grossmann *et al.*, 2003]. In the following, the data of the instruments providing the highest time resolution and located closest to the radical measurements are used, i.e. the Hantzsch data for HCHO.

3. Modeling Approach

3.1. Master Chemical Mechanism

[17] The “master chemical mechanism” (MCM) [Jenkin *et al.*, 1997] was used to calculate the OH, HO_2 , and RO_2 radical concentrations from the photolysis frequencies and trace gases that were measured simultaneously at Pabstthum. The MCM consists of 10 500 chemical reactions involving 17 inorganic and 120 organic species. The chemical mechanism was updated in the following way: the α -pinene oxidation scheme was adopted from the MCM2 model (the MCM2 model is available at <http://www.chem.leeds.ac.uk/Atmospheric/MCM>). The rate coefficients for the reaction of NO_2 with OH, and for the reaction of O_3 with NO were updated according to Sander *et al.* [2000] and

the rate coefficient for the reaction of HO_2 with NO was revised to $9.7 \times 10^{-12} \text{ cm}^3 \text{ s}^{-1}$ based on the measurements made by Bohn and Zetzsch [1997]. The photolysis frequencies which are calculated within MCM from the solar zenith angle were tied to the measurements in the following way: For $J(NO_2)$ and $J(O^1D)$, the values measured by the two filter radiometers were used directly. The other photolysis rates were calculated in the MCM model for clear-sky conditions and then scaled with the ratio of the measured to calculated values of $J(NO_2)$.

[18] The differential equations were integrated with the software package FACSIMILE [Curtis and Sweetenham, 1987]. The model was initialized every 5 min with the measured concentrations of NO, NO_2 , O_3 , CO, PAN, HCHO and higher carbonyl compounds, NMHCs, H_2O , and aerosols, as well as temperature and photolysis frequencies (see Table 1). The initial radical concentrations were taken from the output of the previous 5 min time interval. The time depended concentrations yielded radical concentrations very close to steady state conditions.

3.2. Simple Quasi-Stationary State Model

[19] It is interesting to compare the radical ratios HO_2/OH and RO_2/HO_2 obtained from the MCM with calculations using a simple steady state approach based on the reactions shown in Table 2. We estimate the concentrations of the short-lived radicals species by assuming that the HO_2/OH ratio is determined predominantly by radical recycling reactions. The validity of this assumption is demonstrated in section 5.1 where radical budgets are discussed. In the approximation for RO_2/HO_2 ratio we neglect the secondary production of peroxy radicals due to RO decomposition and isomerisation, as well as the production of RO_2 and HO_2 from ozonolysis, and the loss reactions due to RO_2 self reactions, due to its minor contributions. Adding up the major production and loss terms for both, HO_2 and RO_2 radicals, one obtains for the steady state of HO_2 (14) and RO_2 (15):

$$\begin{aligned}
 -d[HO_2]/dt = 0 = & k_3[OH][CO] + k_4[OH][HCHO] \\
 & + k_{5i}[RO_2][NO] + 2J(HCHO)_{\text{radical}}[HCHO] - k_2[HO_2][NO] \\
 & - k_{10}[RO_2][HO_2] - k_9[HO_2][HO_2]
 \end{aligned} \quad (14)$$

$$\begin{aligned}
 -d[RO_2]/dt = 0 = & k_{6i}[OH][VOC_i] \\
 & + k_{12}[PAN]\{(k_{11}[NO_2]/k_{13}[NO] + 1)\}^{-1} + k_7[OH][CH_4] \\
 & - k_{5i}[RO_2][NO] - k_{10}[RO_2][HO_2]
 \end{aligned} \quad (15)$$

By assuming quasi steady state between $[HO_2]$ and $[RO_2]$, the HO_2/OH ratio can be approximated from reactions (14) and (15):

$$\begin{aligned}
 \frac{[HO_2]}{[OH]} \\
 \approx \frac{(k_{6i}[VOC_i] + k_3[CO] + k_4[HCHO] + k_7[CH_4] + (Q + P)/[OH])}{k_2[NO]}
 \end{aligned} \quad (16)$$

The term $P = k_{12}[PAN]\{(k_{11}[NO_2]/k_{13}[NO]) + 1\}^{-1}$ denotes the production of CH_3O_2 from PAN decomposition

Table 2. List of Chemical Reactions That Have a Major Influence on the Radical Budget of OH, HO₂, and RO₂

Species	Radical Formation	k or J Parameter	Reaction Number
OH	H ₂ O + O(¹ D) → OH + OH	k ₁	(1)
OH	HONO + hν → OH + NO	J _(HONO)	
OH	HO ₂ + NO → OH + NO ₂	k ₂	(2)
HO ₂	HCHO + hν + O ₂ → 2HO ₂ + CO	J _(HCHO)	
HO ₂	OH + CO + O ₂ → HO ₂ + CO ₂	k ₃	(3)
HO ₂	OH + HCHO + O ₂ → HO ₂ + CO + H ₂ O	k ₄	(4)
HO ₂	HO ₂ + NO + O ₂ → HO ₂ + NO ₂ + Carbonyl	k ₅	(5)
RO ₂	OH + VOC + O ₂ → RO ₂ + H ₂ O	k ₆	(6)
RO ₂	OH + CH ₄ + O ₂ → CH ₃ O ₂ + H ₂ O	k ₇	(7)
Species	Radical Destruction	k or J Parameter	
OH	OH + NO ₂ (+M) → HNO ₃ (+M)	k ₈	(8)
OH	OH + VOC + O ₂ → RO ₂ + H ₂ O	k ₆	(6)
OH	OH + CO + O ₂ → HO ₂ + CO ₂	k ₃	(3)
OH	OH + CH ₄ + O ₂ → CH ₃ O ₂ + H ₂ O	k ₇	(7)
OH	OH + HCHO + O ₂ → HO ₂ + CO + H ₂ O	k ₄	(4)
HO ₂	HO ₂ + NO → OH + NO ₂	k ₂	(2)
HO ₂	HO ₂ + HO ₂ → H ₂ O ₂ + O ₂	k ₉	(9)
HO ₂	HO ₂ + RO ₂ → ROOH + O ₂	k ₁₀	(10)
RO ₂	RO ₂ + NO + O ₂ → HO ₂ + Carbonyl + NO ₂	k ₅	(5)
RO ₂	RO ₂ + NO → α RONO ₂	k _{5a}	(5a)
RO ₂	RO ₂ + HO ₂ → ROOH + O ₂	k ₁₀	(10)
<i>PAN Decomposition as Radical Source</i>			
RO ₂	CH ₃ C(O)O ₂ + NO ₂ → PAN	k ₁₁	(11)
RO ₂	PAN → CH ₃ C(O)O ₂ + NO ₂	k ₁₂	(12)
RO ₂	CH ₃ C(O)O ₂ + NO + O ₂ → CH ₃ O ₂ + CO ₂ + NO ₂	k ₁₃	(13)

according to the reactions (11)–(13) (see Table 2). While the total PAN decomposition proceeds at a much larger rate, most of the peroxyacetyl radicals react with NO₂ back to PAN. The term Q denotes the HO₂ production from the radical channel of HCHO photolysis: $Q = 2J(\text{HCHO})_{\text{radical}} \times [\text{HCHO}]$. Photolysis of the higher carbonyl compounds is neglected in (16) as it contributes less than 10% of HCHO to the peroxy radical production at Pabstthum [cf., Grossmann *et al.*, 2003]. The OH concentration on the right side of reaction (16) is taken from the measurements. The term $(Q + P)/[\text{OH}]$ makes a relatively small contribution and is in the range of 10 to 20% of the nominator in reaction (16).

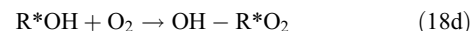
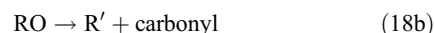
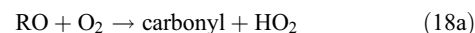
[20] The ratio of RO₂ and HO₂ radicals is then given by reaction (17):

$$\frac{[\text{RO}_{2i}]}{[\text{HO}_2]} \approx \frac{\left(\frac{k_2}{k_5}\right)}{\left\{ \frac{k_3[\text{CO}] + 2J(\text{HCHO})_{\text{radical}}[\text{HCHO}]/[\text{OH}] + k_4[\text{HCHO}]}{(k_{6i}[\text{VOC}] + P)/[\text{OH}]} + 1 \right\}} \quad (17)$$

In this simplified approach, the RO₂/HO₂ ratio depends on the ratio of the rate constants k₂ and k₅ and on the ratio of the production rates for HO₂ and RO₂ radicals which are calculated from the measurements. The average rate coefficient for the reaction of organic peroxy radicals with NO $k_5 \approx 8.6 \times 10^{-12} \text{ cm}^3 \text{ s}^{-1}$ is estimated from the partitioning of the organic peroxy radicals calculated by the MCM (see Figure 3).

[21] Reaction (17) provides a lower limit of the RO₂/HO₂ ratio, because of the neglecting of the secondary production of RO₂ radicals from the decomposition and isomerisation of the RO intermediates. The major pathway for RO radicals in the atmosphere is the reaction with O₂, forming HO₂ and a carbonyl compound [Atkinson, 1997] (18a). There is

substantial evidence from laboratory experiments, that some of the larger RO radicals decompose through a C-C (or C-H) bond splitting (18b) or isomerise via an internal H-shift ((18c) and (18d)) at a rate that is larger than their reaction with O₂ [Atkinson, 1997; Le Bras, 1997]. In the presence of O₂, both processes lead to new peroxy radicals, which again convert NO to NO₂. As a consequence, these reactions increase the ozone formation potential of certain VOCs with respect to the primary OH attack.



Inclusion of the additional RO₂ production by equations (18b)–(18d) would lead to an increase in the steady state concentration of RO₂ and hence of the RO₂/HO₂ ratio. However, the effect is difficult to treat quantitatively, because the effective branching ratios for reactions (18) are not well known. Vice versa, experimental quantification of the RO₂/HO₂ ratio can help to investigate, how many organic peroxy radicals are formed upon the OH-initiated hydrocarbon oxidation.

4. Results

4.1. Diurnal Variation of the Radical Concentrations

[22] The peroxy radical concentrations measured by MIESR during the first two intensive days of the BERLIOZ campaign (20–21 July 1998) are shown in Figure 1. Panel

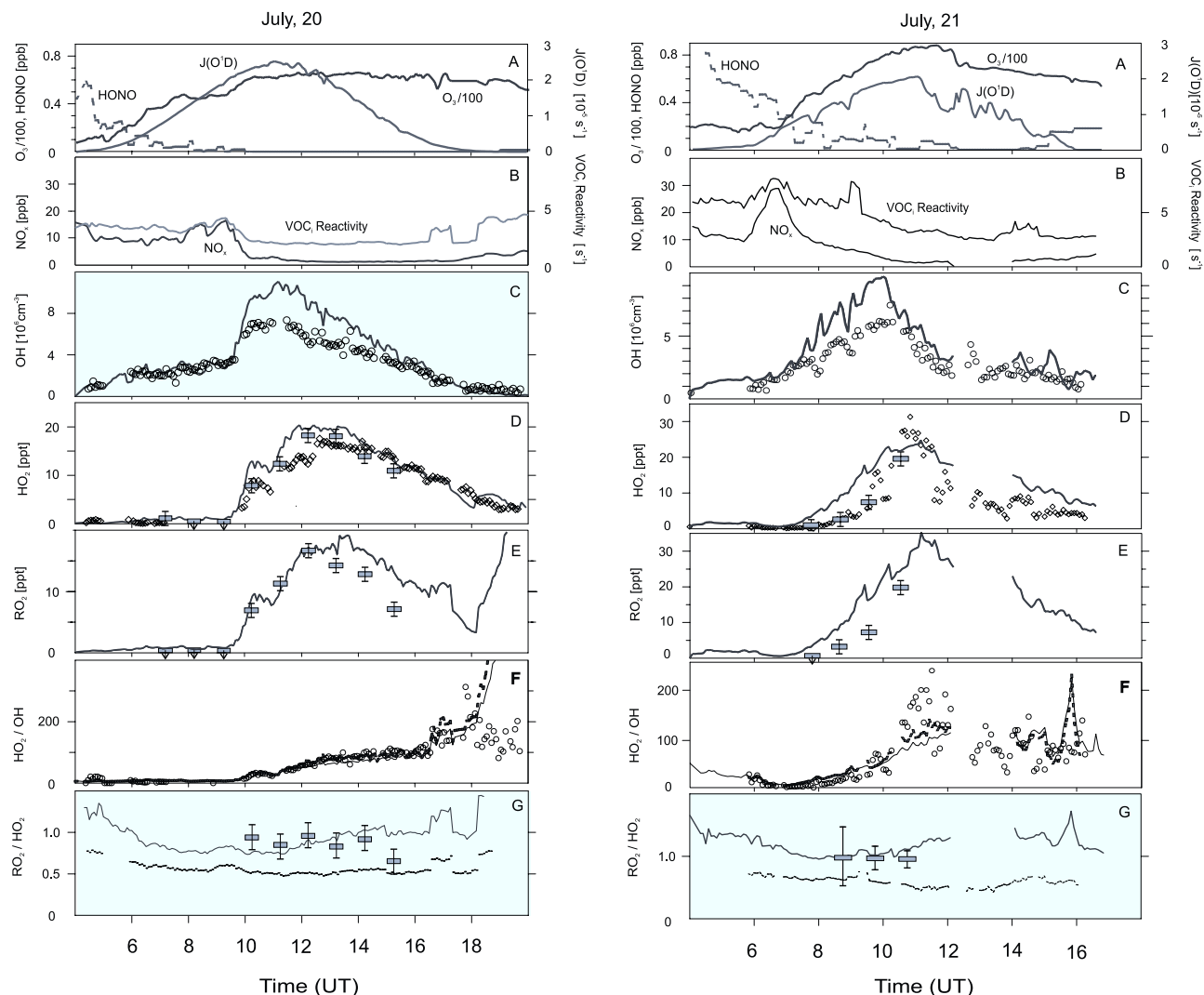


Figure 1. Diurnal variation of the radical concentrations and some additional parameters at Pabstthum on (left) 20 July and (right) 21 July. Panels show the following: A, HONO, O_3 , and $J(O^1D)$; B, NO_x and total VOC reactivity; C, OH concentration measured by LIF (open circles); D, HO_2 measured by LIF (open circles) and by MIESR (rectangles); E, Sum of all organic peroxy radicals (RO_2) measured by MIESR (rectangles); F, HO_2/OH ratio derived from the LIF data; G, RO_2/HO_2 ratio derived from the MIESR data. The MCM results are shown as solid lines in panels C–G. The dotted lines in panels F and G are results of the simplified steady state model (equations 16 and 17).

D gives the concentration of HO_2 and panel E the total concentration of organic peroxy radicals RO_2 . Also shown are the OH and HO_2 concentrations measured by LIF [Holland *et al.*, 2003] in panels C and D, respectively. The upper two panels display some of the measured parameters that are important for the radical budgets, i.e. $J(O^1D)$ [Holland *et al.*, 2003], O_3 and NO_x , [Volz-Thomas *et al.*, 2003b], HONO [Alicke *et al.*, 2003], and the total reactivity of the VOC toward OH, i.e. $\Sigma([VOC_i] \cdot k_i)$ [Konrad *et al.*, 2003]. The lowest panels show the HO_2/OH and RO_2/HO_2 ratios.

[23] The first day was characterized by two different air masses with quite different chemical composition. The transition between the two regimes occurred within 30 min between 9:30 and 10:00 UT as can be seen from the NO_x mixing ratio and VOC reactivity in panel B. The

relatively high- NO_x concentrations (~ 10 ppb) at night are most likely due to local emissions from the surrounding grass lands under a shallow inversion and at low wind speeds [Glaser *et al.*, 2003; Volz-Thomas *et al.*, 2003b]. After break up of the inversion, the city plume of Berlin was encountered at Pabstthum around 9 UT with NO_x mixing ratios of up to 18 ppb. After the air mass change, NO_x dropped to levels below 5 ppb. We shall denote the two regimes as high- NO_x (>10 ppb) and low- NO_x (<10 ppb) in the following.

[24] The peroxy radical concentrations show qualitatively the expected anticorrelation with NO_x . At high- NO_x very low concentrations at or below the detection limit (2ppt) of MIESR are found. After the air mass change, both, HO_2 and RO_2 start to rise and reach mixing ratios of 15–20 ppt around local noon, from where they steadily decrease in the

course of the afternoon. The HO₂ mixing ratio measured by LIF is in very good agreement with the MIESR data, except for the sample collected around 12:00 UT on 20 July. As discussed by [Volz-Thomas *et al.*, 2003b], good agreement is also found for the sum of HO₂ and RO₂ by MIESR with the measurements made by chemical amplification. The OH concentration measured by the LIF instrument peaks about 2 hours earlier than the peroxy radicals. It is noteworthy, that OH is present at fairly large concentrations of $(2-4) \times 10^6 \text{ cm}^{-3}$ during the morning hours when the NO_x mixing ratio is still greater than 10 ppb. The concentration of CH₃C(O)O₂, which can be discriminated from the other peroxy radicals by its spectroscopic signature [Mihelcic *et al.*, 1990], never exceeded the detection limit (2 ppt) of the MIESR. The meteorological situation on 21 July was similar to the day before, until a frontal system with scattered thunderstorms approached the region in the afternoon. In order to prevent damage by lightning strokes, the MIESR measurements were stopped at 11 UT and also the NO_x measurements were switched off for calibrations between 12:20–14:00 (UT). The NO_x mixing ratios in the morning were even higher than on 20 July with maximum values of about 30 ppb. The balloon soundings [Glaser *et al.*, 2003] showed that the vertical extension of the high-NO_x regime was only about 100 m. The strong decrease of NO_x and other anthropogenic pollutants after 7:00 (UT) is mostly explained by the growing height of the mixed layer. The low wind speeds at ground and the wind direction preclude the explanation that the NO_x was advected from Berlin as on the day before [Glaser *et al.*, 2003]. HONO exhibited a quite similar behavior as on the previous day with a maximum in the early morning hours and a strong decay after sunrise due to increasing photolysis. Other than on 20 July, isoprene as an indicator for biogenic emissions from the nearby forest predominated the VOC's reactivity on 21 July.

[25] The diurnal variation of the radical concentrations was quite similar to that on the previous day with HO₂ and RO₂ peaking around local noon and OH somewhat earlier. The maximum concentrations of the peroxy radicals were somewhat larger, whereas the OH concentration was slightly lower than on 20 July. Again, relatively high-OH concentrations were observed during the morning in the presence of large NO_x mixing ratios. The faster decay of all radicals in the afternoon is consistent with the reduced UV radiation flux due to the appearance of clouds when the frontal system approached the measurement site.

4.2. MCM-Model Calculations of Radical Concentrations

[26] The MCM simulations of OH, HO₂ and RO₂ are compared with the measurements in Figure 1. On 20 July, the modeled OH concentrations agree with the measured data within the range of the measurement uncertainty during the time of the plume period between 6:00 and 9:30 (UT). During the transition to less polluted air, the model and the observations both show a sharp rise of OH. After the air mass has changed, the model significantly overpredicts OH by a factor 1.6 until about 16:00 (UT). For HO₂ the model behaves similarly with respect to the observations. During the high-NO_x conditions in the morning, it agrees well with the measurements which are at the detection limit. After the

air mass has changed, the model overpredicts the measurements until about 14:30 (UT). Relative to the MIESR data the overprediction is 50% at 10:15 UT and about 10% at 11:00–13:00 UT. With respect to the LIF measurements, the overprediction is 50% around 12:00 UT. The modeled values of RO₂ agree with the measured data in the morning until 12:00 UT. In the afternoon there is an increasing overprediction by the model reaching a factor 1.7 at 15:00 (UT). The sharp increase in the modeled RO₂ concentration after 18.00 is due to the onset of nighttime chemistry when the production of RO₂ occurs mainly via ozonolysis and NO₃ reactions, as is discussed in detail by [Geyer *et al.*, 2003].

[27] On the next day the model behaves quite similar as on the day before. While good agreement is again observed during early morning in the high-NO_x regime, the model overpredicts the OH, HO₂, and RO₂ concentrations by about a factor of two during the rest of the day. An exception is the temporary agreement of the modeled HO₂ data with the measurements at 11:00 UT. No comparison can be made for RO₂ before 7:45 UT and during the afternoon when corresponding measurements were missing.

[28] The ratio RO₂/HO₂ derived from the MIESR measurements shows little variability on both days and lies mostly in the range between 0.75 and 1.0. Note that the ratio is only shown for data above the detection limit of MIESR, i.e., between 10:00–15:00 UT on 20 July and 9:00–11:00 UT on 21 July. The experimental ratios are reasonably well reproduced by the model (Figure 1), which is a consequence of the fact that the overpredictions for HO₂ and RO₂ cancel to a large extent. In the early morning and late evening the modeled RO₂/HO₂ ratio is higher than during midday and increases to values around 1.3.

[29] The ratio HO₂/OH derived from the LIF measurements on 20 July increases from about 30 in the morning to about 170 at 17:00, with decreasing NO concentrations. The ratio was typically between 10 and 60 under polluted conditions, where NO was in the range of 0.2 to 0.7 ppb, and between 80 and 140 when NO was low (0.1 to 0.2ppb). For 21 July the observed values of the HO₂/OH ratio are much more scattered than on 20 July and reach maximum values of 240 around noon. The ratios observed on 20 July are reproduced within 20% by the model over the whole day until sunset. On 21 July, however, the model results resemble the observations only roughly. The model does not capture the observed variability of HO₂/OH, leading to sporadic positive or negative deviations that exceed a factor of 2. The comparison of modeled and measured radical concentrations is investigated more quantitatively in Figure 2 in form of scatterplots, in which the data are filtered for high-NO_x (>10ppb) and low NO_x (<10ppb) conditions. In case of the peroxy radicals the high-NO_x data are omitted, because the measurements were at the detection limit of the instruments, thus yielding no meaningful correlation with the model. For OH the correlations show a very similar picture on both days. At high-NO_x (upper panels) there is agreement for OH within the experimental uncertainties. We like to note that HONO photolysis plays an important role during this part of the day and that neglecting this OH source would lead to a strong under prediction of OH [cf., Alicke *et al.*, 2003]. In the low NO_x regime (<10ppb), the model overpredicts OH by 50 percent or more on both days, with slightly more scatter on

21 July. For the peroxy radicals the picture at low NO_x is quite different for the two days. A relatively tight linear correlation between modeled and measured HO_2 and RO_2 concentrations with a slope of 1.1–1.2 is observed on 20 July. At lower HO_2 concentrations on 21 July the model overestimated the measurements below 10 ppt and leveled off at higher concentration. Similar behavior is suggested for the RO_2 concentrations. Figure 3 shows the partitioning of the organic peroxy radicals as predicted by MCM for the two days. According to the model, methylperoxy and other aliphatic RO_2 radicals comprise more than 50% of RO_2 . Oxidation of olefins and carbonyl compounds accounts for the remaining fraction. Interestingly, the peroxyacetyl mixing ratio predicted by MCM does not exceed the detection limit of the MIESR (2 ppt), which is in agreement with the measurements. Peroxy radicals from α -pinene oxidation are negligible (<1ppt) on both days. The major difference between the two days is in the fraction of RO_2 originating from isoprene oxidation which are unimportant on 20 July except for the late afternoon. On 21 July, isoprene oxidation accounts for 20–50% of the RO_2 in the MCM over the entire day. This is due to the 5 times larger isoprene mixing

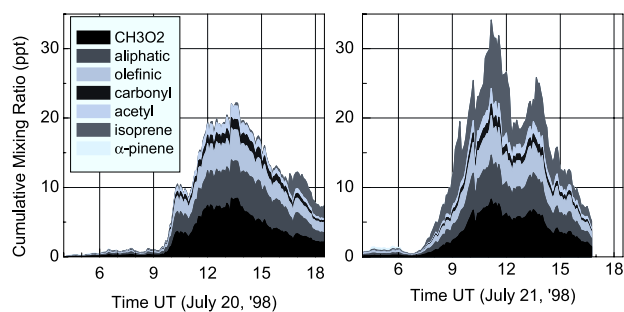


Figure 3. Speciation of RO_2 as calculated by the MCM for (left) 20 July and (right) 21 July. See color version of this figure at back of this issue.

ratio on that day [cf., Konrad *et al.*, 2003]. These authors show that isoprene predominated the total hydrocarbon reactivity on 21 July. A possible suggestion is therefore, that the chemical mechanism implemented in MCM significantly overestimates the peroxy radical production from isoprene oxidation.

5. Discussion

5.1. Radical Budgets

[30] Figures 4a and 4b show the major rates of production and destruction for the different radicals. The reactions that are taken into account are listed in Table 2. The rates were calculated from the measured values of the radicals, trace gases, and the photolysis frequencies, using the same rate coefficients as used in MCM. In general the rates of the recycling reactions which interconvert OH, HO_2 , and RO_2 are much faster than those of the primary production by photolysis of O_3 , HONO, and HCHO, or radical loss by radical-radical recombination. Only during the early morning, the OH is budget strongly influenced by the primary production via photolysis of HONO and O_3 , and destruction by the reaction of OH with NO_2 .

[31] The major OH production term is due to recycling reaction of $\text{HO}_2 + \text{NO}$ (about 70%). The remaining OH production is associated to photolysis of O_3 . During the high- NO_x period, destruction of OH with NO_2 and VOC are about equal, followed by the reaction with CO. In the low NO_x period, reactions with VOC predominate the OH loss. On 21 July the OH destruction rate by VOCs is larger and more variable as on the previous day because of the much higher concentrations of isoprene, most likely due to nearby emissions.

[32] The major HO_2 production term during the low NO_x period is the reaction of RO_2 with NO (about 60%), followed by reactions of OH with CO and CH_2O , as well as photolysis of CH_2O . Photolysis of the higher aldehydes and ketones contributes only a few percent. Destruction of HO_2 is mainly caused by reaction with NO (>80%). For high- NO_x conditions an estimate production and destruction rates is difficult since the measured HO_2 and RO_2 concentrations are below the detection limit.

[33] The peroxy radical budget is dominated by reactions of OH with VOCs (50–70%), followed by production of CH_3O_2 due to PAN decomposition (term P in equation (16)). In the afternoon of 21 July the net PAN decom-

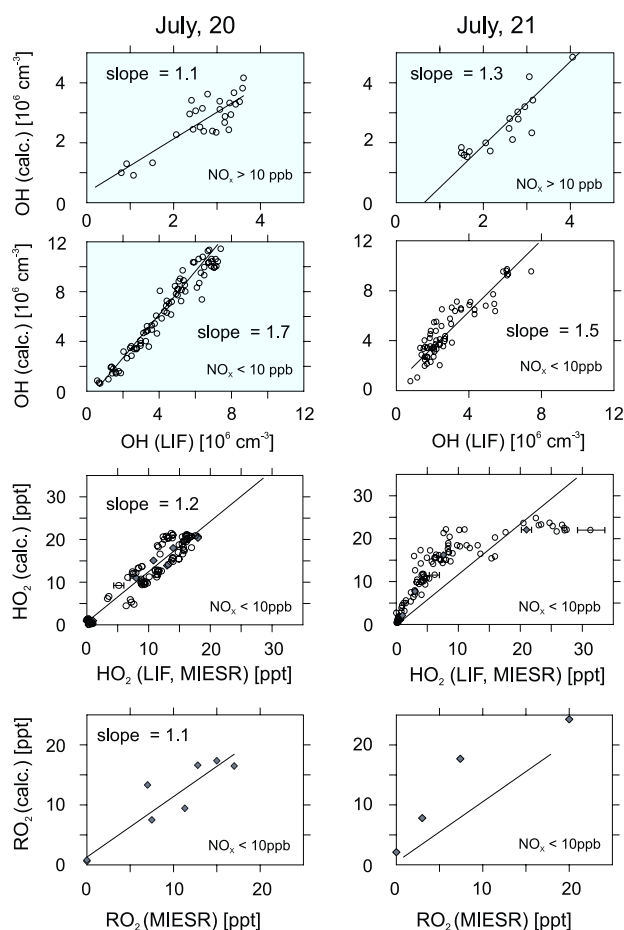


Figure 2. Correlation of calculated (MCM) and measured radical concentrations at Pabstthum for (left) 20 July and (right) 21 July. For OH the high- NO_x and low- NO_x regimes are shown separately. HO_2 and RO_2 for $\text{NO}_x > 10$ ppb are not shown (below detection limits of MIESR and LIF). The solid lines represent linear fits with the indicated slopes.

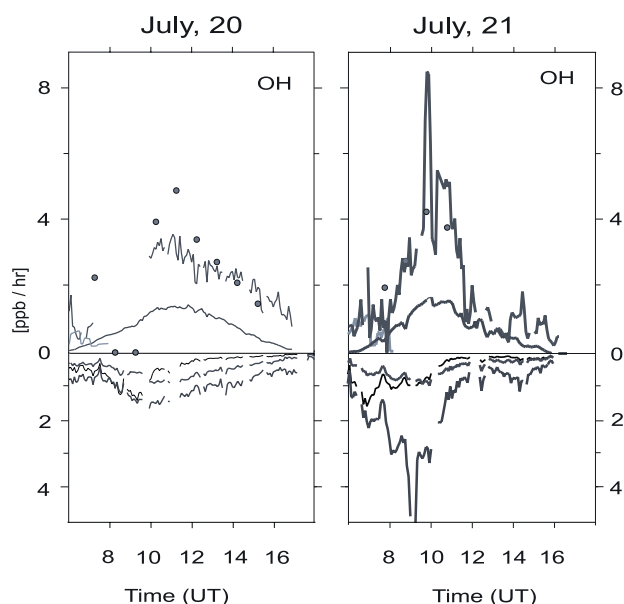


Figure 4a. Major production and destruction rates of OH during (left) 20 July and (right) 21 July calculated from the observations. OH: $k_2 \times \text{HO}_2 \times \text{NO}$ (green line: LIF, gray circles: MIESR); $k_1 \times \text{H}_2\text{O}_2 \times \text{O}(^1\text{D})$ (pink line); $\text{J}(\text{HONO}) \times \text{HONO}$ (yellow line); OH destruction: $k_8 \times \text{OH} \times \text{NO}_2$ (black line); $k_i \times \text{OH} \times \text{VOC}_i$ (red line); $k_3 \times \text{OH} \times \text{CO}$ (brown line). All values are in units of ppb/hr. See color version of this figure at back of this issue.

position equals the production rate by OH + VOC. Destruction of the RO_2 radicals is always dominated by reaction with NO. Reactions of RO_2 radicals with HO_2 comprise a minor contribution, except for the MIESR sample collected around noon on 21 July. The gross destruction rate by reaction of RO_2 with NO reaches 3.5 ppb/hr two times larger than the total RO_2 production rate on 20 July. The RO_2 destruction and production rates on 21 July are reasonably balanced.

[34] The ratios between the total production and destruction rates (P/D) of OH, HO_2 and RO_2 are shown in Figure 5. The data are shown only for the low NO_x regime, where the MIESR measurements were significantly above the detection limit. In case of OH and HO_2 it is found that the total production rates are generally larger than the total destruction rates. The ratio P/D for RO_2 is quite different for the two days. On 20 July the RO_2 production is considerable smaller than destruction, except for the last measured point. On 21 July the ratio P/D varies between 0.7 to 1.7. We note, however, that recycling of RO_2 from RO decomposition and isomerisation is missing in the RO_2 budgets. The contribution is difficult to calculate, as is described in section 3.2. In order to estimate its influence, MCM model results are used to correct the ratio P/D which is then shifted to net production ($1 < \text{P/D} < 2.3$) on both days.

[35] The budget of HO_2 is governed by production from $\text{RO}_2 + \text{NO}$ and destruction by $\text{HO}_2 + \text{NO}$ (Figures 4a and 4b). The destruction and production terms are in balance within 10% on 20 July (Figure 5, middle), whereas production exceeds destruction by up to a factor of 1.6 on 21 July (Figure 5, middle).

[36] OH production exceeds destruction by up to a factors of 2 and 3 on 20 and 21 July, respectively (Figure 5). Production of OH is mainly due to reaction of HO_2 with NO during these periods (Figures 4a and 4b). The two independent HO_2 measurements (LIF and MIESR) agreed within 10%. The accuracy of the NO measurements is estimated to 5% [Volz-Thomas *et al.*, 2003a, 2003b]. Likewise, the parameters used to calculate the OH production from ozone photolysis have accuracies of 5% for O_3 , H_2O and 14% for $\text{J}(\text{O}^1\text{D})$. Therefore, the imbalance is unlikely caused by an overestimation of the production terms, but points to a missing sink for OH. This imbalance in the OH budget is in agreement with the overprediction of OH by the model (Figure 1), which forces a balanced budget.

[37] Overprediction of OH by constrained models was observed in a number of studies [McKeen *et al.*, 1997; Hauglustaine *et al.*, 1999; George *et al.*, 1999; Carslaw *et al.*, 1999] and several mechanisms, i.e. missing VOC reactivity and heterogeneous losses of OH and/or HO_2 on particles, were discussed as possible solutions for the observed discrepancy. In order to balance the OH production rates for 20 and 21 July an additional reactivity of 250–300 ppt isoprene equivalents is required ($0.6\text{--}0.75 \text{ s}^{-1}$). Such an additional amount of VOC reactivity would lead to a corresponding increase in the production of RO_2 . Likewise, including such a reactivity in the MCM leads to an

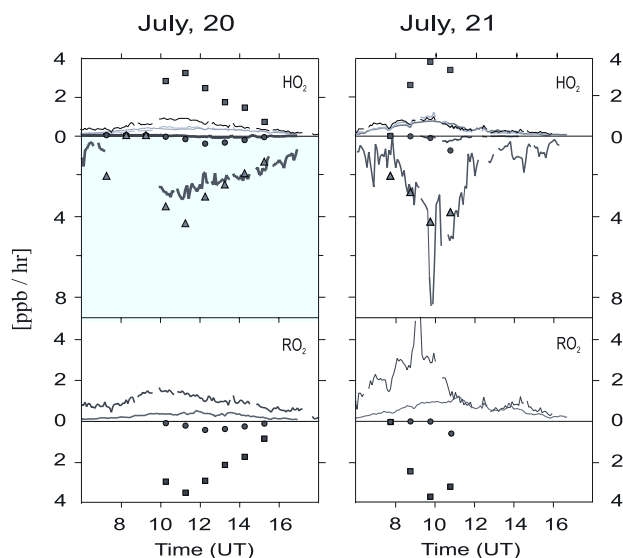


Figure 4b. Major production and destruction rates of HO_2 and RO_2 during (left) 20 July and (right) 21 July calculated from the observations. (top) HO_2 : $k_5 \times \text{HO}_2 \times \text{NO}$ (blue squares); $k_3 \times \text{OH} \times \text{CO}$ (black line); $k_4 \times \text{OH} \times \text{HCHO}$ (yellow line); $2\text{J}(\text{HCHO})_{\text{radical}} \times \text{HCHO}$ (gray line); HO_2 destruction: $k_2 \times \text{HO}_2 \times \text{NO}$ (green line, LIF) and pink triangles (MIESR); $k_{10} \times \text{HO}_2 \times \text{RO}_2$ (brown circles, MIESR); $k_9 \times \text{HO}_2 \times \text{HO}_2$ (red line, LIF). (bottom) RO_2 : $k_i \times \text{OH} \times \text{VOC}_i + k_7 \times \text{OH} \times \text{CH}_4$ (red line); CH_3O_2 production from PAN decomposition (blue line); RO_2 Destruction: $k_5 \times \text{RO}_2 \times \text{NO}$ (green squares); $k_{10} \times \text{HO}_2 \times \text{RO}_2$ (brown circles, MIESR). All values are in units of ppb/hr. See color version of this figure at back of this issue.

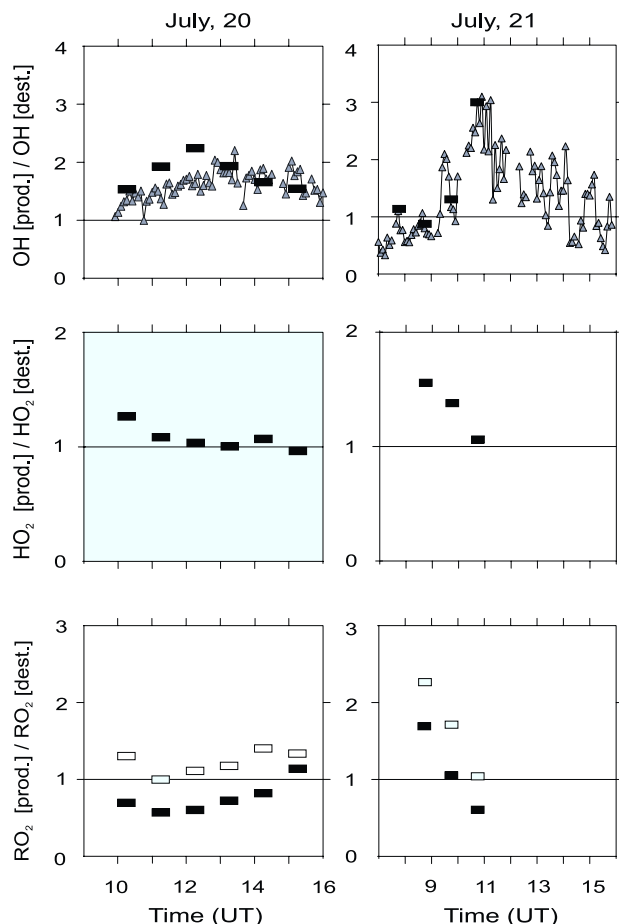


Figure 5. Diurnal variation of the ratio of total production and destruction rates for the different radicals on (left) 20 July and (right) 21 July for (top) OH (triangles from LIF data, rectangles from MIESR data), (middle) HO₂ (from MIESR data), and (bottom) RO₂ (from MIESR data, solid rectangles without RO chemistry, open rectangles including RO chemistry).

overestimation of the RO₂ mixing ratio by almost a factor of two. As a consequence, one would have to invoke VOCs or some other unknown species that do not produce RO₂ radicals upon oxidation by OH.

[38] The potential influence of heterogeneous radical losses on the budgets of OH, HO₂ and RO₂ was investigated in several studies [Cantrell *et al.*, 1996; Saylor, 1997; Jacob, 2000; Kanaya *et al.*, 2000]. At first approximation, the loss rates can be estimated from the specific surface area (*A*) of the particles and an uptake coefficient equation [cf., Carslaw *et al.*, 1997]:

$$k = \gamma \times (RT/2\pi)^{0.5} \times A \quad (19)$$

The specific surface area *A* (m²/m³) was measured on both days with maximum values of 220×10^{-6} (m²/m³) on 20 July and 450×10^{-6} (m²/m³) on 21 July (Alicke, personal communication, 1999). The uptake coefficient for HO₂ was found to be $\gamma > 0.025$ on ice and $\gamma > 0.07$ on sulphuric acid (28%). For CH₃C(O)O₂, γ was found to be even smaller, ~ 0.004 on water and ~ 0.001 on sulphuric acid (51%)

[DeMore *et al.*, 1997]. With these values for γ , there was little difference (<5%) in the modeled OH, HO₂ and RO₂ concentrations. In order to match the measured radical concentrations on 20 July an uptake coefficient of 0.5 is needed for both HO₂ and RO₂, similar to the value used by Cantrell *et al.* [1996]. For 21 July, reasonable agreement within 20% is achieved for the OH concentration. The modeled HO₂ and RO₂ concentration are reduced by 20%, thus the overprediction remains high and is not resolved by including heterogeneous losses.

[39] A significant difference between the two days is the larger role of PAN in the RO₂ budget on 21 July because of slightly higher concentrations and the higher temperature. We have investigated the sensitivity of MCM to PAN by setting its concentration to zero. This extreme case scenario reduced the models overprediction of RO₂ on 21 July by about 20% and had practically no influence on the RO₂ concentrations on 20 July. Therefore errors in the rate coefficients for PAN decomposition or formation are unlikely the reason for the discrepancy in the RO₂ budget on 21 July.

5.2. HO₂/OH Ratios

[40] The discussion of the radical budgets has shown that the production and destruction of OH, HO₂, and RO₂ is dominated by the reactions which convert these radicals into each other. Accordingly, the HO₂/OH ratio would be expected to be controlled by the interconversion reactions as estimated by the simple steady state expression in reaction (16). The result of this simple model is plotted in panel F of Figure 1 (left and right) (dotted lines) and can be compared to the full MCM model calculation (solid lines). On both days the simple equation and the complex MCM model yield very similar results. The model predictions are in general good agreement with the observed HO₂/OH ratio on 20 July, which suggests that the interconversion reactions for OH and HO₂ are sufficiently well understood on this day. This is apparently not the case on the other day, 21 July, where the MCM and the simple model both fail to reproduce the observed variability of HO₂/OH. The variance is most likely due to strong fluctuations in the concentrations of isoprene and other biogenic VOCs that are advected from the nearby forest. These variations are not resolved by the gas chromatographic measurements which represent 20 min averages [Konrad *et al.*, 2003]. The apparent discrepancy between model and radical concentrations thus highlights the need for time resolved VOC measurements in situations, where close by sources cannot be ruled out. Quite different results have been reported recently by Tan *et al.* [2001] for an environment, where much higher isoprene concentrations (several ppb) were present. In the PROPHET campaign they measured OH and HO₂ inside a deciduous forest at low NO_x (<1ppb) for low NO_x conditions. For the modeled HO₂ they found good agreement with the observations, whereas the model was generally unable to match the measured OH, with the observations 2.7 times greater than the model on average. As a result, the modeled HO₂/OH ratios were 2.5–4 times higher than the measured ratios, indicating that the cycling between OH and HO₂ was poorly described by the model [Tan *et al.*, 2001]. The results from PROPHET and BERLIOZ suggest that biogenic VOC possibly have a complex

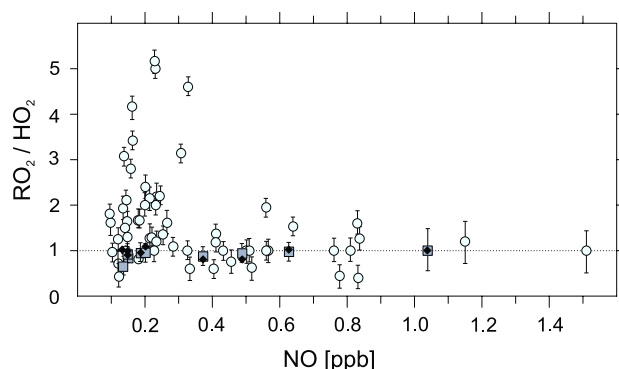


Figure 6. Ratios of RO₂ to hydroperoxy radicals for different NO concentrations measured by MIESR during field campaigns in 1989, 1990, 1992 and 1994 at Schauinsland (open circles) and during BERLIOZ (measurements shaded square, and solid diamonds, MCM calculations). The data represent half hourly values and are filtered for $J(\text{NO}_2) \geq 0.004 \text{ s}^{-1}$.

influence on the tropospheric photochemical system which is not yet well understood.

5.3. RO₂/HO₂ Ratios

[41] Figure 6 shows the dependence of RO₂/HO₂ as measured at Pabstthum with those obtained during several previous campaigns at Schauinsland in the Black Forest, Germany, as a function of the prevailing NO mixing ratio. Moreover, the MCM model predictions for Pabstthum are included in the Figure 6. The data are selected for $J(\text{NO}_2) > 0.004 \text{ s}^{-1}$. The measurements at Schauinsland show RO₂/HO₂ ratios of up to 5 at low NO concentrations. The ratio declines rapidly at $\text{NO} > 0.3 \text{ ppb}$ to a value of around 1. Note that the RO₂ and HO₂ concentrations are near to or below the detection limit of the MIESR for $\text{NO} > 1.6 \text{ ppb}$. For the conditions at Pabstthum the MCM predicts RO₂/HO₂ ratios of about one, almost independent of the NO mixing ratio between 0.05 and 10 ppb in excellent agreement with the MIESR data. The simple steady state model (reaction (17)) predicts much lower ratios since it does not account for RO₂ recycling from the decomposition and isomerisation of the alkoxy radicals. Obviously, MCM describes most of these processes correctly.

[42] *Stevens et al.* [1997] discuss RO₂/HO₂ ratios for the TOPHE campaign at Idaho Hill which were derived from the combination of CA data (RO₂ + HO₂) and HO₂ measurements by LIF. They find ratios of about 4–5 under clean conditions ($\text{NO} < 0.2 \text{ ppb}$) in good agreement with the data from Schauinsland. The model used by *Stevens et al.* [1997] predicts a RO₂/HO₂ ratio of about one, in agreement with the MCM results for Pabstthum. It seems therefore that the current chemical mechanisms miss an important process that leads to higher RO₂/HO₂ ratios at low NO_x concentrations.

[43] Under polluted conditions ($\text{NO} > 1.5 \text{ ppb}$), however, these authors find RO₂/HO₂ ratios of 12–14, much higher than what is observed at either Pabstthum or Schauinsland. The very large RO₂/HO₂ ratios observed by *Stevens et al.*

[1997] in the presence of 1.6–1.9 ppb NO are almost exclusively due the very high RO₂ + HO₂ concentrations (5–11 ppt) measured by the chemical amplifier [*Cantrell et al.*, 1997]. The coexistence of such high RO₂ and NO mixing ratios is impossible to explain with our current understanding of the radical chemistry and disagrees with MIESR measurements from several campaigns. *Stevens et al.* [1997] discuss two possible solutions for the discrepancy between the model and their measurements: A 10 times smaller value for the rate coefficient of reaction (5) RO₂ + NO and a factor of 5 larger rate coefficients for the recombination reactions of RO₂ with HO₂. The major improvement comes from reaction (5), since the peroxy radical recombination has little influence on the RO₂/HO₂ ratio except for conditions where NO_x is extremely low. The VOC reactivity at Idaho Hill under high-NO_x conditions is dominated by isoprene of ($\sim 2.1 \text{ s}^{-1}$), followed by ethene ($\sim 1 \text{ s}^{-1}$), propene ($\sim 1 \text{ s}^{-1}$) and HCHO of ($\sim 0.65 \text{ s}^{-1}$). It is difficult to believe that the RO₂ radicals, produced by oxidation of these hydrocarbons, would react 10 times more slowly than recommended by *Atkinson* [1997]. For example, the measured rate constants for the hydroxyalkyl peroxy radicals HOCH₂CH₂O₂ ($9 \times 10^{-12} \text{ cm}^3 \text{ s}^{-1}$ [*Le Bras*, 1997]) and OHC(CH₃)₂CCH₂O₂ ($4.9 \times 10^{-12} \text{ cm}^3 \text{ s}^{-1}$ [*Wallington et al.*, 1997]) are within a factor 2 of the recommendation for the corresponding alkyl peroxy radicals [*Atkinson*, 2000]. Also, a 10 times smaller rate coefficient for RO₂ + NO would lead to even larger discrepancies for the NO/NO₂ ratio derived from the direct measurements and from the photostationary state (PSS) (see discussion by *Volz-Thomas et al.* [2003b]).

[44] Aerosol formation during the photooxidation of biogenic VOC would be another possible reason for decreasing the HO₂ production from the initial RO₂ radicals. The yield of aerosol formation was measured by *Pandis et al.* [1991] for several compounds and ranged from negligible in the case of isoprene, 30–40% for β -pinene and limonene, to 100% for trans-caryophyllene. *Hoffmann et al.* [1997] note, however, that ozonolysis is responsible for most of the aerosol formation in the oxidation of the biogenic monoterpenes and sesquiterpenes in their studies.

6. Ozone Production Rates

[45] It is generally assumed [cf., *Volz et al.*, 1988; *Ridley et al.*, 1992; *Kleinman et al.*, 1995; *Frost et al.*, 1998] that the photochemical ozone formation rate $P(\text{O}_3)$ is approximately determined by the rate at which NO is oxidized to NO₂ by reaction with peroxy radicals (HO₂ and RO₂ (see reactions (2) and (5) in Table 2). In this approach, the minor pathway of the higher organic peroxy radicals that leads to formation of organic nitrates and the loss of NO₂ by reaction with OH are neglected. In this respect reaction (20) represents an upper limit for $P(\text{O}_3)$.

$$P(\text{O}_3) = [\text{NO}] \times \{k_2[\text{HO}_2] + \sum(k_{si}[\text{RO}_2]_i)\} \\ \cong [\text{NO}] \times \{k_2[\text{HO}_2] + k_5[\text{RO}_2]\} \quad (20)$$

The ozone production rate data for Pabstthum on 20 and 21 July are shown in Figure 7 as a function of NO,

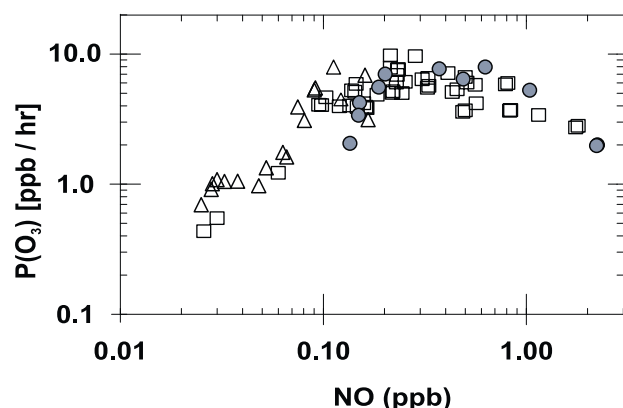


Figure 7. Comparison of the ozone production rate calculated from the measured radical concentrations (reaction 20) during the BERLIOZ campaign (solid circles) to those calculated from the MIESR measurements made during several campaigns at Schauinsland (summer 1989, 1990, 1992 and 1994, squares) and in August 1993 at Izana, Tenerife (triangles). The simultaneously observed NO mixing ratio is used as abscissa. The data represent half hourly values and are filtered for $J(\text{NO}_2) \geq 0.004 \text{ s}^{-1}$.

denoted by the filled circles. Note that both scales are logarithmic. Also shown are values for $P(\text{O}_3)$ obtained from earlier campaigns at Schauinsland (1989, 1990, 1992, 1994) and Tenerife [1995]. In each case, $P(\text{O}_3)$ was calculated from the peroxy radical concentrations measured by MIESR and the average NO concentration in the MIESR sampling period data and selected for $J(\text{NO}_2) > 0.004 \text{ s}^{-1}$. An average rate coefficient of $k_5 = 8.6 \times 10^{-12} \text{ cm}^3 \text{ s}^{-1}$ for the reaction of organic peroxy radicals with NO was estimated from the evaluation of the different rate coefficients by [Atkinson, 1997] and the partitioning of the radicals provided by MCM. For HO_2 , the rate coefficient by Bohn and Zetzsch [1997] was used.

[46] The observed O_3 production rate at Pabstthum shows a strong increase with the NO mixing ratio between 0.1 and 0.3 ppb, where $P(\text{O}_3)$ reaches a maximum value of 8 ppb/hr. The dependence of $P(\text{O}_3)$ on NO at concentrations below 0.3 ppb has a positive slope of 53 ppb/hr per ppb NO. Above 0.5 ppb NO the rate of ozone production decreases linearly with a slope of 3.3 (ppb/hr)/(ppb NO). This behavior can be compared with the $P(\text{O}_3)$ data obtained from Schauinsland and Tenerife. The corresponding peroxy radical and NO measurements at Schauinsland were made in May 1989, August 1990, September 1992, and August 1994. [Mihelcic and Volz-Thomas, 1994]. At Izana, Tenerife, measurements were performed during the OCTA (Oxidizing Capacity of the Tropospheric Atmosphere) intensive in August 1993 [Schultz, 1995; Fischer et al., 1998; Zenker et al., 1998]. The $P(\text{O}_3)$ data from these campaigns show a similar behavior as was observed at Pabstthum. A strong, nearly linear increase with the NO mixing ratio is found between 0.02 and 0.25 ppb, where $P(\text{O}_3)$ reaches a maximum value of 10 ppb/hr. At NO values above 0.3 ppb, $P(\text{O}_3)$ decreases slowly with further increasing NO concen-

trations. The linear increase below 0.3 ppb NO has a slope that is about 30% smaller than observed during the BERLIOZ campaign, whereas the decreasing trend of $P(\text{O}_3)$ above 0.5 ppb NO is very similar to what is found at Pabstthum. The scatter of the $P(\text{O}_3)$ data over the whole range of the NO mixing ratios is relatively low, suggesting some similarity of the ozone production for the three different places in Europe.

[47] A similar dependence of the ozone production rate on the NO concentration was calculated using a steady photochemical model by Kleinman et al. [2000]. The calculated $P(\text{O}_3)$ show a maximum value of up to 11 ppb/hr, and shows nearly linear dependence of $P(\text{O}_3)$ on NO, at NO concentrations below 0.25 ppb [Kleinman et al., 2000]. Frost et al. [1998] investigated the dependence of the ozone production on NO_x . They found a slope for low $[\text{NO}_x]$ (< 2 ppb) of about 6 [ppb/hr]/[ppb NO_x]. This is in the same order of magnitude as our observation of about 7 [ppb/hr]/[ppb NO_x]. Nevertheless, their maximum value of ozone production, calculated from $[\text{RO}_x]$ measurements by chemical amplifier techniques [Cantrell et al., 1992, 1993] was about a factor of 4 higher than ours, because of the higher isoprene concentrations during the ROSE campaign.

7. Summary and Conclusions

[48] In this paper we have analyzed radical measurements of OH, HO_2 , and RO_2 , which were obtained in a rural environment near the city of Berlin on two intensive days (20 and 21 July) in summer 1998. On both days the atmospheric conditions were characterized by high- NO_x levels (10–30 ppb) in the morning, followed by a transition to relatively low NO_x concentrations, which were about an order of magnitude smaller and persisted until the evening. The two days were distinctly different with respect to the chemical composition of the VOC. On 21 July high isoprene levels were encountered, indicating the advection of biogenic VOC emissions from a nearby mixed forest.

[49] The concentrations and diurnal variations of the radicals showed a similar behavior for the two days. Relatively high OH concentrations of $(2\text{--}4) \times 10^6 \text{ cm}^{-3}$ were observed at high NO_x in the morning, while HO_2 and RO_2 remained near the detection limit ($\sim 1\text{--}2$ ppt) for these conditions. At low NO_x the concentrations of OH reached $(6\text{--}8) \times 10^6 \text{ cm}^{-3}$ at local noon, while HO_2 and RO_2 reached 20–30 ppt, each. The HO_2/OH ratio varied from 10 to 240, with higher values observed at low NO_x . The RO_2/HO_2 ratio was quite constant and showed values between ~ 0.75 and 1.0 during daytime.

[50] The observed radical concentrations and their ratios have been compared with model simulations using a Master Chemical Mechanism (MCM) that was constrained by measured values of a comprehensive suite of trace gases and photolysis frequencies. Main features of the diurnal radical cycles are reasonably reproduced by the model, for example the strong increase of the radicals during the transition from high- to low- NO_x conditions. At high NO_x the concentration of OH is well described by the model with only slight overpredictions ($\sim 10\%$ on July 20, $\sim 30\%$ on 21 July). The analysis of the radical budget has shown that in this case OH was strongly

influenced by the primary production via photolysis of HONO and O₃, and the destruction of OH via the reaction with NO₂. For the peroxy radicals the model predictions agree with the observed low concentration values. A precise comparison is not possible in this case, because the peroxy radical measurements were at the detection limit.

[51] At low NO_x the model generally tends to overestimate all radicals (OH, HO₂, RO₂) by varying amounts. OH is overpredicted by a factor of 1.5–1.7 on the two days. In case of the peroxy radicals the overprediction on July 21 is larger than on the previous day, reaching a factor of 2 or more, when isoprene concentrations were high. In general the ratios HO₂/OH and RO₂/HO₂ are much better reproduced by the model than the individual measurements. On 20 July the modeled ratios agree within 20% with the observations. On the following day, the model fails to reproduce the large variability of the experimental HO₂/OH ratio. The observed variance on this day was likely due to fluctuations of isoprene and other biogenic VOC from nearby sources, which were not resolved by the gas chromatographic measurements and therefore not appropriately considered in the model.

[52] The analysis of the radical budgets has been performed almost completely on the basis of field observations. The analysis shows that the recycling reactions, which interconvert OH, HO₂, and RO₂, dominate the chemical turnover and the ratios of the radicals under the conditions of BERLIOZ at NO_x levels below 10 ppb. Thus, the agreement of the modeled and observed ratios HO₂/OH and RO₂/HO₂ demonstrates that the radical recycling is reasonably well described by the model, at least for the conditions on 20 July. The measured radical ratios have also been compared with the result of a simple steady state model, which only accounts for cycling reactions of the radicals. The simple model reproduces the observed HO₂/OH ratio as well as the complex MCM model, but underestimates the RO₂/HO₂ ratio by a factor 1.2–1.8. This factor reflects the efficiency of the neglected RO₂ recycling from the decomposition and isomerisation of the alkoxy radicals. Apparently the MCM model accounts for most of these processes correctly.

[53] The general model overprediction of the radicals at low NO_x is consistent with the empirical radical budgets, which show that the total production rates are larger than the total destruction rates for this condition. Since there is no experimental evidence that the production rates are determined too high, the budgets can only be balanced by additional radical sinks, which do not recycle OH, HO₂, or RO₂. No process could be identified that would quantitatively explain this kind of radical loss. Missing VOC in the model could be responsible for the overprediction of OH, but would not explain the overprediction of HO₂ and RO₂. Loss of OH by the reaction with HX (X = Br, I) would lower the radical concentrations, but unreasonably high HX concentrations in the order of one ppb would be required to explain the observed OH. Heterogeneous loss processes on aerosol can be ruled out as an unlikely explanation, since an uptake coefficient of about 0.5 would be needed, which is much larger than laboratory measurements suggest.

[54] The measured peroxy radicals have been used to estimate the local ozone production rate during BERLIOZ. The production rate was found to increase linearly with NO below 0.3 ppb NO and to decrease above 0.5 ppb NO. The maximum rate of 8 ppb/hr was encountered at 0.3 ppb NO. A similar behavior of the ozone production rate has been derived from previous peroxy radical measurements at Schauinsland in the Black Forest and Tenerife, where the maximum O₃ production never exceeded values of 10 ppb/hr.

[55] In conclusion, the comprehensive data set of radical measurements at BERLIOZ has given interesting insight into the photochemistry of the lower troposphere and has confirmed important aspects in the tropospheric master chemical mechanism. There remain, however, significant deficiencies in the model, such as the model overprediction of the radicals, for which no conclusive explanation has been found. The role of biogenic VOC as a sink of OH and source of peroxy radicals requires more attention. In this respect, the development of faster VOC measurement techniques with a time resolution of about a few minutes would support greatly future field studies of the fast radical photochemistry.

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- G. K. Moortgat, Max-Planck-Institut für Chemie, Division of Atmospheric Chemistry, Postfach 3060, D-55020 Mainz, Germany.

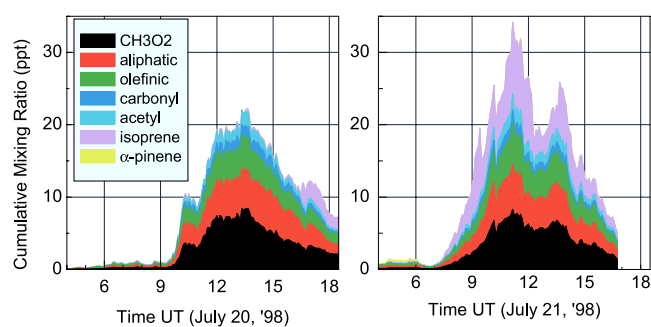


Figure 3. Speciation of RO_2 as calculated by the MCM for (left) 20 July and (right) 21 July.

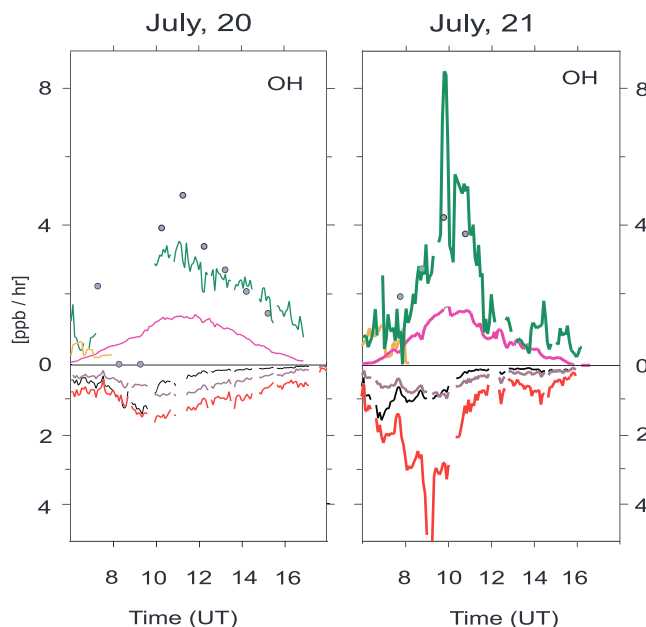


Figure 4a. Major production and destruction rates of OH during (left) 20 July and (right) 21 July calculated from the observations. OH: $k_2 \times \text{HO}_2 \times \text{NO}$ (green line: LIF, gray circles: MIESR); $k_1 \times \text{H}_2\text{O} \times \text{O}(^1\text{D})$ (pink line); $J(\text{HONO}) \times \text{HONO}$ (yellow line); OH destruction: $k_8 \times \text{OH} \times \text{NO}_2$ (black line); $k_1 \times \text{OH} \times \text{VOC}_i$ (red line); $k_3 \times \text{OH} \times \text{CO}$ (brown line). All values are in units of ppb/hr.

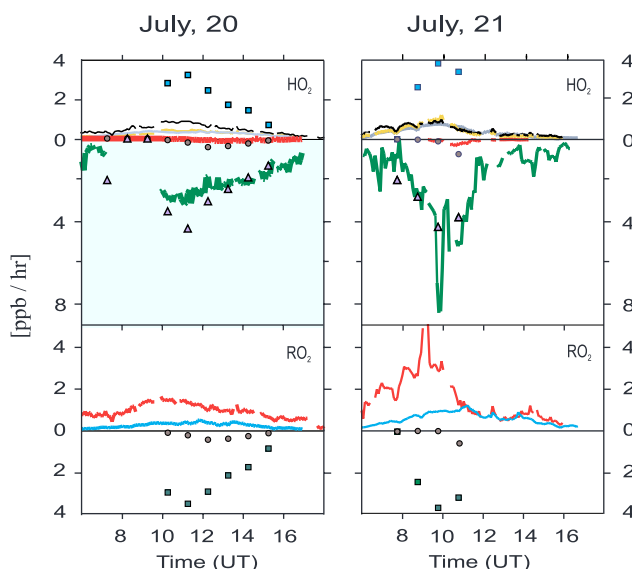


Figure 4b. Major production and destruction rates of HO_2 and RO_2 during (left) 20 July and (right) 21 July calculated from the observations. (top) HO_2 : $k_5 \times \text{RO}_2 \times \text{NO}$ (blue squares); $k_3 \times \text{OH} \times \text{CO}$ (black line); $k_4 \times \text{OH} \times \text{HCHO}$ (yellow line); $2J(\text{HCHO})_{\text{radical}} \times \text{HCHO}$ (gray line); HO_2 destruction: $k_2 \times \text{HO}_2 \times \text{NO}$ (green line, LIF) and pink triangles (MIESR); $k_{10} \times \text{HO}_2 \times \text{RO}_2$ (brown circles, MIESR); $k_9 \times \text{HO}_2 \times \text{HO}_2$ (red line, LIF). (bottom) RO_2 : $k_1 \times \text{OH} \times \text{VOC}_i + k_7 \times \text{OH} \times \text{CH}_4$ (red line); CH_3O_2 production from PAN decomposition (blue line); RO_2 Destruction: $k_5 \times \text{RO}_2 \times \text{NO}$ (green squares); $k_{10} \times \text{HO}_2 \times \text{RO}_2$ (brown circles, MIESR). All values are in units of ppb/hr.