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Article

Characterization of Total OH Reactivity in a Rapeseed Field: Results from the COV³ER Experiment in April 2017

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Abstract: Croplands remain poorly studied ecosystems in terms of total hydroxyl radical (OH) reactivity, especially when compared to forests. As part of the COV³ER project, total OH reactivity (ROH), defined as the total loss rate of OH due to its reaction with reactive species in the atmosphere, was characterized in a rapeseed field (Grignon, France) during the blooming season in April 2017. Measurements were performed in a dynamic chamber as well as in ambient air using the Comparative Reactivity Method (CRM). Complementary measurements of organic (including a proton transfer reaction quadrupole ion–time of flight mass spectrometry, PTRQI-ToFMS) and inorganic compounds were also performed in order to calculate the expected OH reactivity and evaluate the missing fraction. Measured ROH varied diurnally in the dynamic chamber ($mROH_{\text{chamber}}$) with maxima around 20 to 30 s^{-1} at midday and minima during dark hours, following the variability of the enclosed branch $VOCs_{\text{rapeseed}}$, which is light- and temperature-dependent. Oxygenated VOCs were the major compounds emitted by the rapeseed crop. However, in terms of contribution to OH reactivity, isoprene accounted for 40% during the daytime, followed by acetaldehyde (21%) and monoterpenes (18%). The comparison between $mROH_{\text{chamber}}$ and calculated ROH ($cROH_{\text{chamber}}$) exhibited little or no difference during dark hours, whereas a maximum difference appeared around midday, highlighting a significant missing fraction (46% on average during daytime) mainly related to biogenic temperature- and/or light-dependent emissions.

Keywords: OH reactivity; biogenic VOCs; CRM-PTRMS; PTRQI-ToFMS; crops

1. Introduction

Up to 10^4 – 10^5 of different organic species have been identified so far in the Earth's atmosphere and this figure may only represent a small fraction of the total number actually present [1]. They arise

from anthropogenic and natural sources, the latter emissions being dominant (90%) on the global scale [2]. Once emitted in the atmosphere, volatile organic compounds (VOCs) are transformed by photochemical processes, leading to the formation of harmful secondary pollutants such as ozone (O_3) and secondary organic aerosols (SOAs), which have serious impact on human health and the environment. The major depletion route of atmospheric VOCs is via reaction with OH radicals, representing the main tropospheric oxidant during the daytime [3].

Natural sources of VOCs include forests, agricultural lands, grasslands, shrubs, biomass burning and oceans [4]. They are involved in growth, reproduction and defense mechanisms of the plant and they play an important role in allelopathic interactions, defined as chemically induced interactions between plants or fungi [5,6]. It has been reported that biogenic VOCs' (BVOCs) chemical composition and emission rates are linked to internal factors in the plant like genetics and biochemical processes. They can be affected by abiotic and biotic external factors such as temperature, radiation, water availability and interactions with other living species [7,8]. Biogenic VOCs exhibit a large chemical diversity. They include terpenoids, which are dominant in the global atmosphere (isoprene, monoterpenes, sesquiterpenes, etc.), oxygenated VOCs (OVOCs such as methanol, acetone, acetaldehyde, formaldehyde, etc.) and other classes such as sulfur- and nitrogen-containing compounds, organic halides, alkanes and alkenes [9]. All these compounds represent a strong potential sink for atmospheric oxidants and hence to the hydroxyl radical.

One way to characterize the overall sink of OH in a given environment and to evaluate the importance of atmospheric photochemistry is to measure total OH reactivity. Total OH reactivity is defined as the total loss rate of OH, which is the inverse of OH lifetime (s^{-1}). It represents the sum of the reaction frequencies with OH of all chemicals, calculated as the product of each compound concentration by the compound rate coefficient of reaction with OH [10]. Measurements of total OH reactivity have been performed using Laser Induced Fluorescence systems (pump-and-probe or flow reactor) and the Comparative Reactivity Method (CRM) using a Proton-Transfer-Reaction Mass Spectrometer (PTR-MS) as a detector or a fast Gas-Chromatographic/Photo-Ionization Detector, each system exhibiting advantages and limitations [11]. Total OH reactivity proved to be an effective parameter for quantifying the contribution of VOCs to atmospheric chemistry in various environments. However, the comparison between the measured and the calculated OH reactivity from measured species often revealed differences, showing that there are still some unidentified or unquantified OH sinks [12].

Since the 1960s, the focus was mainly made on BVOC emissions from forests, and little interest was shown for BVOC emissions from agriculture [9]. Karl et al. (2009) [13] introduced a new plant-specific emission inventory where they showed that, on the Pan-Europe scale, agriculture contributes up to 27% to total BVOC emissions. These emissions are dominated by OVOCs (65%), followed by monoterpenes (26%) and isoprene (7%). However, they also reported a lack of experimental data on terpenes, sesquiterpenes and oxygenated VOCs, especially from crops, leading to higher uncertainties on BVOC emission estimates from agriculture included in inventories. Similarly, in terms of OH reactivity, most of the studies conducted in natural ecosystems were in forested areas. Only two took place in suburban areas, near agricultural lands [14,15] where different observations were reported. Ren et al. (2005) [14] found a mean OH reactivity of $6.1 s^{-1}$, with little diurnal variation, in an agricultural research farm in Pennsylvania, US, between May and June. Lou et al. (2010) [15] described a diurnal cycle of OH reactivity in a regional background site surrounded by farmlands in the Pearl River Delta, China, during summer, with a mean maximum of $50 s^{-1}$ at dawn dominated by anthropogenic emissions and a mean minimum of $20 s^{-1}$ at noon, more influenced by local biogenic emissions. These differences highlighted the need to better characterize the OH reactivity in rural environments, in which BVOC emissions can add a significant contribution of photochemical pollution due to the surrounding abiotic VOC emissions [16]. In particular, assessment of the OH reactivity at the branch scale, using dynamic chambers, would be a relevant approach to improve our understanding of the photochemical impacts of agriculture. So far, two studies have investigated OH reactivity using dynamic branch

enclosures [17,18]. The first one, conducted by Kim et al. (2011) [17] on four different tree species (red oak, white pine, beech and red maple) and the second by Nolscher et al. (2013) [18] on Norway spruce trees studying total OH reactivity emission rates (TOHRE). No study was found on OH reactivity from a crop dynamic chamber.

Agricultural lands occupy around 37% of land use worldwide [19]. With the increase in global human population (10.2 billion by the year 2100), the increased demand for food and nutrition would require an additional ≈ 2 billion hectares of cropland [20]. Furthermore, in order to meet the increasing need of energy in the modern society, biofuels from plants appear to be an important mitigation strategy for the impact of fossil fuels on the environment [21]. Thus, a new era of agriculture is encouraging the cultivation of biofuel-producing crops such as soybean, corn and rapeseed. Oilseed rape or rapeseed (*Brassica napus*) is one of the main oil and protein-producing plants grown in Europe [22,23]. It represents up to 69% of oleaginous crops and 14% of total agricultural area of cereals-, oil- and protein-producing crops in France [24]. The rapeseed flowering period is considered as a period of high photochemical activity, especially on clear days where the light yellow surface of the crop reflects the sunlight, increasing the radiation density and leading to the emission of various amounts of BVOCs [25]. Some studies have reported that oilseed growing can also be responsible for allergic effects especially at the mid-flowering stage of development [23,26]. Previous studies investigated the variety and amount of BVOCs emitted by the spring rapeseed, showing that OVOCs and monoterpenes were the main emitted compounds. Gonzaga Gomez et al. (2019) [27] found methanol to be the most emitted compound, whereas McEwan and Macfarlane Smith (1998) [23] reported high levels of monoterpenes from spring rapeseed, with limonene exhibiting the highest concentration, followed by β -myrcene and sabinene. In light of all these observations, a better characterization of BVOCs and their contribution to OH reactivity in croplands, among which are rapeseed fields, is of great importance.

This study aims to investigate BVOCs and their total OH reactivity on a rapeseed crop level in a typical agricultural ecosystem, based on CRM/PTR-MS. A dynamic chamber was installed in April 2017 on a healthy, initially non-flowered rapeseed plant to perform measurements during the blooming period. In parallel, VOCs were also sampled in the incoming and the outgoing chamber air using a proton transfer reaction–quadrupole ion–time of flight–mass spectrometry (PTRQI-ToFMS). An expected OH reactivity was calculated from PTRQI-ToFMS data and other complementary measurements of organic and inorganic compounds. Thereby, it was possible to estimate the amount of a potential missing OH reactivity and to discuss its possible sources on a diel basis.

2. Experiments

2.1. Field Site Description

As part of the ADEME-COV³ER project (Biogenic Volatile Organic Compounds: Emissions by managed ecosystems), a field campaign was conducted from the 7th to the 25th of April 2017. The ICOS FR-Gri measurement site (Figure 1) is a 19 ha field belonging to the AgroParisTech experimental farm of Grignon (48°51' N, 1°58' E), located about 30 km west from Paris, France. The usual crop rotation at the field site is winter wheat, silage maize, winter barley, with only one former occasional occurrence of rapeseed in 2013. From the south, the east, the north-west and the south-west, the site is surrounded by roads with substantial traffic (5000 to 15,000 vehicles per day, 2010 counts, Statistiques du département des Yvelines pour 2010). From the north, the south and the east, other agricultural lands surround the site. The small (1100 inhabitants, Insee 2015) village of Grignon is located around 700 m west of the measurement site, and an animal farm (more than 200 cows and 1000 sheep) is situated 400 m to the south-west. The main wind directions are north-west during clear days and south-west during cloudy and rainy days [28–30]. Rapeseed was sown on 16 August 2016, with a density of 460 000 plants/ha. The canopy height was around 134 cm.

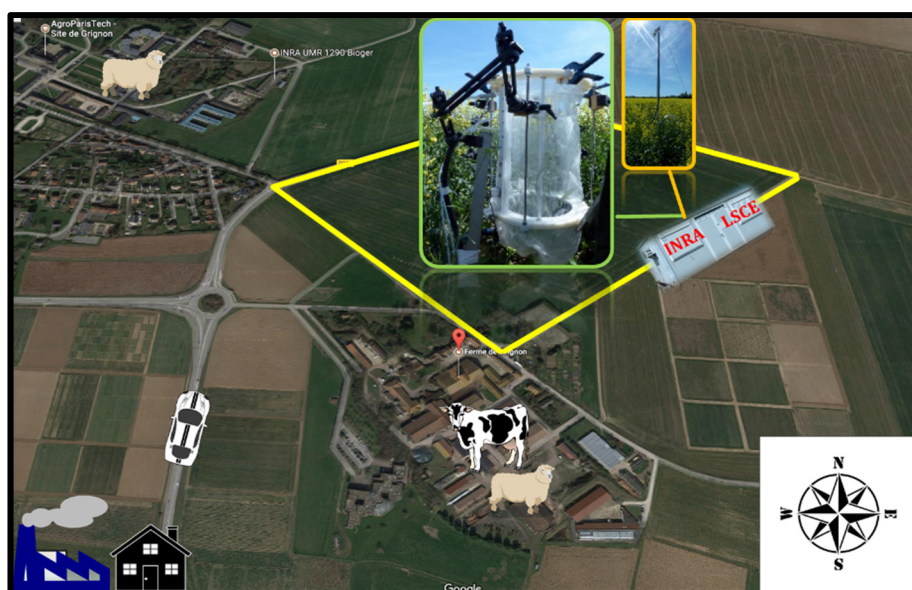


Figure 1. Satellite image of Grignon farm, the surrounding area and the chamber used for this study.

2.2. Plant Dynamic Chamber and Air Sampling

In order to determine the chemical nature and the concentration of BVOCs emitted from a rapeseed plant as well as their OH reactivity, a young, healthy, non-flowered rapeseed plant (*Brassica napus*) was selected for the installation of an enclosure system (a dynamic chamber) from which the measurements were made. The enclosure system was a 60 L polytetrafluoroethylene (PTFE) cylindrical frame bounded by a sealed 50 μm thick PTFE film to which ambient air was introduced at 15 L min^{-1} using a PTFE-coated diaphragm pump, leading to a residence time of about 4 min. Ambient air was chosen instead of zero air, in order to measure BVOC and OH reactivity at the crop level, under in situ conditions, without altering potential plant interaction with ambient air species. Furthermore, the introduction of ambient air is useful to track not only plant emissions, but also the deposition of compounds present in the atmosphere. A rapid mixing of the chamber air and a slight overpressure were ensured using a PTFE fan. Temperature (T $^{\circ}\text{C}$), photosynthetic active radiation (PAR $\mu\text{mol m}^{-2} \text{ s}^{-1}$), relative humidity (RH %), carbon dioxide (CO_2 $\mu\text{mol mol}^{-1}$) and water vapor (H_2O mmol mol^{-1}) were continuously monitored inside the chamber. For more details about the enclosure system refer to Genard-Zielinski et al. (2015) [31].

2.3. Meteorological Measurements

Measurements of meteorological variables were carried out at the FR-Gri ICOS site, according to the ICOS standard protocols [32]. They include two profiles of air temperature and relative humidity (measurement heights at 1, 2.7 and 5 m) and wind speed/direction (measurement heights at 0.5, 1, 2, 3 and 5 m), radiation measurements (short- and long-wave incoming and reflected radiation, photosynthetically active radiation, albedo), soil temperature and humidity profiles from the surface down to 90 cm, and soil surface heat flux. The data were acquired continuously with dataloggers (Campbell Inc., USA) at rates from 20 s to 1 min, quality-checked and post-processed to get half-hourly averages. All sensors were regularly maintained and calibrated to ensure good quality data.

2.4. Instrumentation

2.4.1. Comparative Reactivity Method (CRM) and Instrument Performance

Total OH reactivity measurements were performed using the CRM, first developed and described by Sinha et al. (2008) [33]. This technique relies on quantifying the competition between a reference

molecule (pyrrole C₄H₅N) and ambient reactive species to react with in situ generated OH radicals. The whole experiment can be summarized in three main analytical steps: pyrrole, which is not normally present in ambient air, is introduced with dry zero air and dry nitrogen, in a glass reactor. The latter is equipped with a UV mercury lamp and connected to a detector which in our case is a proton transfer reaction–quadrupole–mass spectrometer (PTR-MS). During this first step, no OH radicals are generated due to dry conditions in the system. The pyrrole concentration is monitored at the protonated *m/z* 68 as C1 which corresponds to the initial concentration of pyrrole available in the reactor after potential photolysis. Then, dry N₂ is humidified and dry zero air is replaced by humid zero air (generated from ambient air going through a catalytic converter) in order to generate OH radicals by H₂O photolysis under the 184.9 nm wavelength emitted by the lamp. Pyrrole concentration at this stage is C2, lower than C1, due to its reaction with OH. The third and final step consists of introducing ambient air instead of humid zero air. The various species present in ambient air compete with pyrrole for the available OH, leading to an increase in pyrrole concentration measured as C3. Measurements of C1, C2 and C3 allow determining the OH reactivity of the sampled air in a quantitative manner where the difference between C2 and C3 is dependent on the amount and reactivity of VOC present in the sampled air. Total OH reactivity is derived from Equation (1), where it is assumed that pyrrole reacts with OH following the pseudo-first-order kinetical regime i.e., [Pyrrole] >> [OH]:

$$R_{\text{OH}} = \frac{(C3 - C2)}{(C1 - C3)} kp C1 \quad (1)$$

with *kp* the reaction rate of pyrrole with OH ($1.2 \times 10^{-10} \text{ cm}^3 \text{ molecule}^{-1} \text{ s}^{-1}$ [34]).

However, three main measurement artifacts can occur while operating the CRM, for which corrections should be applied on OH reactivity raw values: (1) changes in the relative humidity within the setup between zero air (C2) and ambient air (C3), leading to different OH levels, (2) recycling of OH in the presence of nitrogen oxide (NO) due to the NO + HO₂ reaction and (3) deviation from pseudo-first-order kinetics. In some CRM systems, corrections for potential NO₂ and/or O₃ artifacts are also considered [35,36]. An additional correction is applied due to the dilution of ambient air inside the reactor by N₂ and pyrrole flows.

CRM/PTR-MS System Characterization

In order to assess the performance of the instrument and to apply adequate corrections on OH reactivity raw data, a set of tests have been performed throughout the field campaign (see S1). The PTR-MS (Ionicon Analytik GmbH, Innsbruck, Austria) was calibrated at the beginning and the end of the campaign showing a good stability in dry and wet conditions. The photolysis rate $\frac{(C0-C1)}{C0}$ (with C0 being the pyrrole level in dry conditions before switching on the UV mercury lamp) was determined to be around 6% since pyrrole can absorb at some of the wavelengths emitted by the UV mercury lamp. On average, C1 was 70.5 ± 0.3 ppbv for the measurement period presented herein.

Even though a catalytic converter was used to generate humid zero air from ambient air, small differences in humidity are still observed between ambient and zero air. These changes can influence the level of OH radicals between C2 and C3, leading to an artifact in the C2 measurement. Therefore, six humidity tests were done, measuring the change in C2 ($\Delta C2$) with the change of RH, tracked using the variation in the *m/z* 37-to-*m/z* 19 ratio ($\Delta [m/z \text{ 37-to-}m/z \text{ 19 ratios}]$) (with *m/z* 37 being H₃O⁺·H₂O the first water cluster and *m/z* 19 being H₃O⁺, the number of primary ions and their ratio being proportional to the absolute humidity inside the reactor). These tests, performed on incoming and outgoing chamber air were in good agreement, showing a linear relationship between $\Delta C2$ (ppbv) and $\Delta (m/z \text{ 37-to-}m/z \text{ 19 ratio})$ with a slope (mean humidity correction factor) of -119.68 ± 3.2 and $R^2 = 0.84$. The application of this factor on C2 (Equation (1)) gives a corrected C2 noted C2*.

As mentioned before, OH radicals are artificially generated inside the reactor using the photolysis of water vapor. This reaction also results in hydrogen atoms that can be associated with oxygen molecules present in air leading to the formation of HO₂. In the presence of high levels of NO in

sampled air, HO₂ radicals can be rapidly converted into OH radicals resulting in different levels of OH between C2 and C3 and leading to an artifact on C3 measurements. Therefore, tests were performed by introducing different levels of NO (1.2 to 19.6 ppbv) inside the reactor while sampling humid zero air. When the NO level in the system increases, the pyrrole mixing ratio in C3 decreases. The variation of C3 ($\Delta C3$) was thus monitored as the difference between an expected C3 and the measured C3, where the expected C3 is calculated using C1, C2 and an expected OH reactivity due to NO. The relationship between $\Delta C3$ and the NO mixing ratio is not linear. It follows a quadratic regression (S1.b), $\Delta C3 = a [\text{NO}]^2 + b[\text{NO}]$, where [NO] is in ppbv with the following parameters fitted during this pre-experimental test $a = -0.0013$ and $b = 0.3214$ (see also Michoud et al., 2015 [35]). This correction leads to corrected C3 noted C3*. We note that this correction was minor most of the time as the NO mixing ratio was low during the campaign (0.7 ± 0.5 ppbv on average during the considered period except for one peak recorded on the morning of the 21st of April 2017 when the NO mixing ratio reached 46 ppbv around 06:00 UTC (local time = UTC + 2)).

Regarding nitrogen dioxide (NO₂), no tests have been performed on site, since NO₂ is not expected to lead to the formation of secondary OH inside the reactor. However, its conversion into NO showed that it can cause an artifact [35]. During the intercomparison of OH reactivity instruments over the SAPHIR chamber [11], our CRM/PTR-MS suffered from an artifact from high (≥ 10 ppbv) NO₂ concentration, even if this effect was to a lower extent when compared to other CRM instruments. The absolute change in OH reactivity values was about $+1.9 \text{ s}^{-1}$ in the presence of 10 ppbv of NO₂. This correction was only considered in order to estimate the weight of the NO₂ interference on OH reactivity measurements. As NO₂ was on average 12 ppbv during the campaign, this could have led to an underestimation of about 2 s^{-1} of the corresponding OH reactivity.

Based on previous experiments, no dependency was seen for LSCE-CRM regarding O₃ [11]. Therefore, no correction was applied.

OH reactivity values are determined based on Equation (1). This equation assumes that the instrument is operated under pseudo-first-order conditions (i.e., [Pyrrole] \gg [OH]), which is not the case with current CRM instruments. To assess this correction, injections of known concentrations of propane and isoprene were performed during the field campaign. These two gas standards represent extreme cases since propane is a medium reactive VOC in the atmosphere ($k_{(\text{propane-OH})} = 1.09 \times 10^{-12} \text{ cm}^3 \text{ molecules}^{-1} \text{ s}^{-1}$), while isoprene is one of the most reactive ($k_{(\text{isoprene-OH})} = 1.0 \times 10^{-10} \text{ cm}^3 \text{ molecules}^{-1} \text{ s}^{-1}$). The measured OH reactivity obtained from these tests is then compared with the expected OH reactivity, which allows extracting a correction factor that depends on the pyrrole-to-OH ratio [37]. Therefore, standard OH reactivity experiments were conducted at various pyrrole-to-OH ratios ranging between 1.7 and 2.7, which is the range observed most of the time during the COV³ER campaign, leading to a correction factor $F = 1.59x^2 - 7.22x + 9.46$ with $x = \text{pyrrole/OH}$. Finally, correction on reactivity values due to dilution was around 1.44 during the campaign (D).

Taking all mentioned corrections into account, final OH reactivity was determined following Equation (2):

$$R_{\text{OH}} = \frac{(C3^* - C2^*)}{(C1 - C3^*)} \text{kp C1 F D} \quad (2)$$

The limit of detection was about 3 s^{-1} (3σ), based on the calculations described by Michoud et al. (2015) [35] and overall uncertainties have been estimated at 35% (1σ) for the measured OH reactivity by our CRM/PTR-MS [38].

Ambient Air Sampling

The sampling lines were shared with the PTRQI-ToFMS. In total, 8 $\frac{1}{4}$ " internal diameter PFA 20 m lines were sampled at a flow rate of $\sim 7 \text{ L min}^{-1}$. Among these 8 lines, the line from the rapeseed enclosure and the line sampling ambient air at 270 cm height were subsampled to the 4 port-valve of the CRM system, with a flow rate around 1 L min^{-1} , through a $\frac{1}{8}$ " internal diameter PFA line, using two pumps: the first one is a PTFE-coated pump located upstream of the reactor and the other one is

from the Gas Calibration Unit (GCU) used to generate humid zero air from ambient air (Figure S2). All Teflon lines were heated to 50 °C.

Since the PTRQi-ToFMS was used during this campaign to measure gradients, eddy covariance fluxes and the air out of the dynamic chambers, the measurement sequence was programmed as following: 20 min of eddy covariance measurements followed by 5 min of gradient measurement, 3 min in the chamber out and 2 min of background.

2.4.2. Proton Transfer Reaction Quadrupole ion-Time of Flight Mass Spectrometer (PTRQi-ToFMS)

Instrument Setup

A PTRQi-ToFMS (Ionicon Analytik GmbH, Innsbruck, Austria) was used for continuous online measurements of VOC mixing ratios at 10 Hz. The analyzer was described in detail previously ([27,39,40]) and the setup used in this study is detailed in Gonzaga Gomez et al. (2019) [27]. VOCs are protonated inside a reaction tube, then injected into a quadrupole ion guide and electromagnetic lenses that focus the ions into the time-of-flight mass spectrometer (ToF, ToFWerk, Switzerland) which separates the ions by inertia prior to their detection on a multi-channel-plate (MCP) coupled to a time-to-digital converter (Burle Industries Inc., Lancaster, PA, USA). The drift tube pressure was tuned to 4 ± 0.0001 mbar, drift temperature was 80 ± 0.06 °C, and drift voltage E was 995 ± 0.03 V. These conditions ensured an E/N ratio (where E is the electric field strength and N the gas density) of 132 ± 0.03 Td (1 townsend = 10^{-17} V cm²), which was chosen as a good compromise to avoid fragmentation and limit clustering [41,42]. The number of detection channels was set to 240,000 and the mass spectrum spanned 15 to 530 m/z . The extraction rate of ions in the ToF was set to 40 μ s (25 kHz). A total of 2500 extracted spectra were averaged prior to recording at 10 Hz. The proton transfer reaction is effective for compounds having a proton affinity higher than that of water (691.7 kJ mol⁻¹), which means that the major atmospheric compounds are not protonated (N_2 , O_2 , CO_2 , CH_4 , N_2O) allowing for detection of trace compounds.

Acquisition and Pre-processing of PTRQi-ToFMS Data

Data acquisition was performed with homemade Labview[®] software which logged a list of ion peaks integrated online by the TOFDAQ software (TOFWERK, SW). The desired ion peak list was set-up at the start of the experiment based on measured air sampled at the site, analyzed by a homemade Labview peak analyzer. Every 5 min, the data were stored on the hard drive, and a mass calibration was performed with the automatic TOFDAQ software feature using ions m/z 21.0221 ($H_3^{18}O^+$) and m/z 59.049 (acetone $C_3H_6OH^+$), and a 2-s moving average. Mass calibration was quality-checked online by computing the peak mass of m/z 21.022 and m/z 59.049 after calibration and was found to be stable throughout most of the experiment. Unstable mass calibration periods were discarded at the computation stage. The channel's number of the 16-way valve embedded in the PTRQi-TOFMS was also recorded. The synchronized data were pre-processed with another Labview program that computed the mean and standard deviation of the cps at the entry and exit of the chamber as well as at reference level (2.7 m) and in the canopy at 1.34 m. In this pre-processing step, the first 15 s and the last 5 s of a recording were removed prior to computing cps statistics to avoid unstable drift pressures that occurred after each valve switching.

PTRQi-ToFMS Calibration

In order to account for the sensitivity changes, the PTRQi-ToFMS was regularly calibrated. We used a standard cylinder containing 102 ppbv of benzene, 104 ppbv of toluene, 130 ppbv of ethylbenzene and 336 ppbv of xylene (122 ppbv Ortho, 121 ppbv Meta, 123 ppbv Para; BTEX, Messer). Gas from this cylinder was diluted with synthetic air (alphagaz 1, Air Liquide, FR) using two fluorinert-coated mass flow controllers (Bronkhorst), to generate concentrations from 0 to 50 ppbv of BTEX. The calibration was done almost every week. The zero check was performed every 30 min by passing through a hydrocarbon filter (Supelco ref 22445-12 or Restek ref 21991) for 2 min and keeping the last 30 s.

The zero was withdrawn from the normalized cps every 30 min as shown in Equation (S3.1). The sensitivity S_i was calculated as the slope with zero intercept of the regression between mixing ratio in equation (S3.1) assuming $S_i = 1$ and the known mixing ratio. The procedure was applied for the three compounds cited above for the calibration. We found that the sensitivity was 2 ± 0.15 . Based on an intercomparison between our PTRQI-ToFMS and a PTR-MS intercompared within the ACTRIS network [43] we adjusted methanol sensitivity to $S_{\text{methanol}} = 14$ as explained in Gonzaga Gomez et al. (2019) [27]. Except for methanol, a single sensitivity value was applied for all masses, since the protonation reaction constant was assumed equal for all masses.

2.4.3. Gas Chromatography-Flame Ionization Detector (GC- FID)

A gas chromatograph equipped with a Flame Ionization Detector (airmoVOC C₂–C₆, Chromatotec, Saint Antoine, France) was used to measure hydrocarbons in the C₂–C₆ fraction. The sampling was done for 10 min with a flow rate of 11 sccm via a stainless steel inlet, followed by a 20 min analysis. After going through a Nafion dryer, ambient air arrives to a preconcentration unit cooled down to -8 °C, filled with Carboxen, Carboxen B and Carboxen C. A thermodesorption at 220 °C follows and the sample is injected into a metal capillary column (Porous Layer Open Tubular Column PLOT, Al₂O₃/KCl; 0.53 mm inner diameter and 25 m length, Varian Inc, USA). Calibrations were performed frequently using a certified standard VOC mixture (National Physical Laboratory, UK). The overall uncertainty was estimated to be 15% (1 σ) [44].

2.4.4. NO_x Concentration Measurements

Nitrogen oxides (NO_x) concentrations in ambient air were measured with a NO-NO₂-NO_x chemiluminescence analyzer (Model 42-CTL, Thermo Environnement, Mègetec, France). Data were acquired continuously at 1 Hz using a datalogger (Model CR10, Campbell Inc., USA). The NO-NO₂-NO_x instrument was calibrated in the laboratory before being installed on the field, and the span was regularly checked during the campaign (about once every two weeks). The inlet was placed at a height of 2.5 m, on the side of the FR-Gri field.

2.4.5. CO and CH₄ Measurements

A 48i-TL, thermo-Environnement was used to measure carbon monoxide (CO) by the absorption of light in the infrared range. A zero was done every 4 h, and the calibration was checked several times throughout the campaign, by the injection of a pre-calibrated air bottle (≈ 200 ppbv).

Regarding methane (CH₄), a G2201-i Picarro based on Cavity Ring Down Spectroscopy (CRDS), was deployed. CRDS uses a single frequency laser diode to measure specific gas-phase molecules which scatter and absorb light in the near infrared absorption spectrum. By measuring the height of absorption peaks, the concentrations can be determined. More details on the instrument and setup used in this campaign can be found in Assan et al. (2017) [45].

All deployed instruments are summarized in Table 1 and Figure 2.

Table 1. Instruments deployed in the campaign and used for data analysis.

Technique	Measurement	Time Resolution
CRM	Total OH Reactivity (ROH)	10 min
PTRQI-ToFMS	VOCs Formaldehyde, Methanol, Acetaldehyde, Butanol, Acetone, Acetic acid, Isoprene, MethylEthylKetone (MEK), Hexadiene/ GLV, Styrene, Monoterpenes	0.1 s

Table 1. Cont.

Technique	Measurement	Time Resolution
GC-FID	C ₂ –C ₆ HCs Alkanes: Ethane, Propane, i-butane, n-butane, i-pentane, n-pentane, 2-methylpentane, n-hexane Alkenes: Ethene, Propene, Trans-2-butene, 1-butene, i-butene, Cis-2-butene, 1,3-butadiene, Trans-2-pentene, 1-pentene Alkynes: Acetylene	30 min
	NO-NO ₂ -NO _x chemiluminescence analyzer	1 min
G2201-i Picarro	CH ₄	1 min
48i-TL, thermo-Environnement	CO	1 min

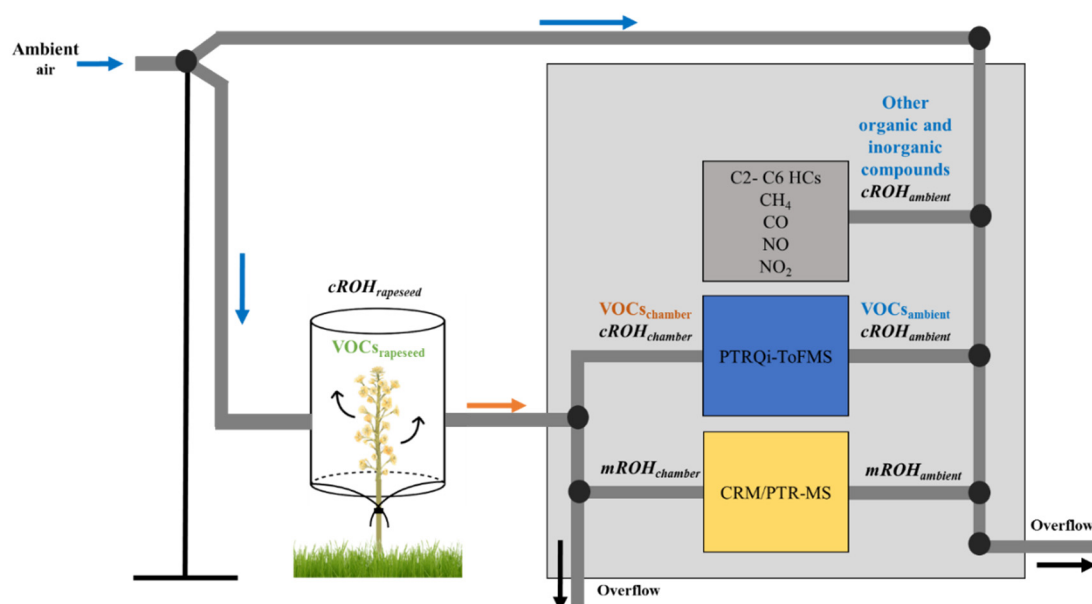


Figure 2. Schematic representation of the system used for the measurements in the incoming and the outgoing chamber air. $mROH_{ambient}$ and $mROH_{chamber}$ stand for measured OH reactivity in the incoming and the outgoing air, respectively. $cROH_{ambient}$ and $cROH_{chamber}$ represent the calculated OH reactivity in the incoming and the outgoing chamber air, respectively. $cROH_{rapeseed}$ is calculated using $VOCs_{rapeseed}$, determined from $VOCs_{chamber} - VOCs_{ambient}$ (both measured with the PTRQI-ToFMS).

2.5. Calculation of OH Reactivity

Based on the available trace gases observations, mentioned in Section 2.4 an expected OH reactivity (calculated OH reactivity, $cROH$) could be determined using the following equation:

$$cROH = \sum cROH_i = \sum k_{i-OH} X_i \quad (3)$$

where $cROH_i$ is the calculated OH reactivity of a compound i , X_i is its concentration and k_{i-OH} is its respective rate coefficient of the reaction with the hydroxyl radical OH.

While $cROH_{ambient}$ was calculated considering $VOCs_{ambient}$ (PTRQI-ToFMS), light (C₂–C₆) hydrocarbons, NO_x, CO and CH₄ measured in ambient air (Figure 2), $cROH_{chamber}$ was determined using $VOCs_{chamber}$ (PTRQI-ToFMS) measured in the outgoing chamber air and the organic and inorganic species measured in ambient air, assuming that inorganic compounds and light hydrocarbons were not emitted nor deposited in the rapeseed chamber. This assumption was made since these compounds'

variabilities in ambient air (S6) showed no dependency on temperature nor PAR, demonstrating that they were not directly linked to biogenic emissions.

As reported in Gonzaga Gomez et al. (2019) [27], the PTRQi-ToFMS deployed during the COV³ER campaign detected 1078 ions (fragments and isotopes included) with a mass-to-charge ratio (m/z) ranging from 31 to 500. For rapeseed, around 420 ions exceeded the set threshold (signal/ noise ratio > 3), from which 24 masses were selected as the main contributors to rapeseed BVOCs emission fluxes, based on their mean net flux. This list of 24 masses was considered first, including m/z 31.018, m/z 33.034, m/z 41.038, m/z 43.018, m/z 45.033, m/z 57.069, m/z 59.048, m/z 61.028, m/z 69.070, m/z 79.063, m/z 81.069, m/z 83.084, m/z 90.946, m/z 91.053, m/z 93.035, m/z 105.069, m/z 108.956, m/z 125.957, m/z 137.131, m/z 153.052, m/z 190.962, m/z 192.960, m/z 223.061 and m/z 225.043. However, when it came to consider VOC_{sambient} and VOC_{schamber} in cROH_{ambient} and cROH_{chamber}, respectively, this list was shortened to 12 masses, with known chemical nature and established rate coefficients (k) for the reaction with OH. It is important to note that, in our case, total monoterpenes are represented by the sum of m/z 81.068 and m/z 137.129, since m/z 81.068 was mainly a monoterpene fragment. Isoprene, detected at m/z 69.070 also exhibited a fragment at m/z 41.038, therefore, the corresponding signal was corrected for the potential fragmentation with a factor of 1.3. Formaldehyde could be detected and quantified at m/z 31.018. Indeed, Warneke et al. (2011) [46] reported that PTR-MS can be used for formaldehyde measurements, even though HCHO detection by PTR-MS can be strongly dependent on humidity and can be less sensitive than for other compounds. The 12 selected PTRQi-ToFMS masses as well as the other organic and inorganic measured trace gases are listed in Table 2, together with their respective rate constants for the reaction with OH used for calculations. It is worth noting that the OH reactivity of monoterpenes was calculated as the product of the concentration of total monoterpenes (m/z 137 and m/z 81) and the rate constant of the reaction of limonene with OH. This assumption was based on previous studies, where limonene was reported to be the dominant monoterpene emitted by *Brassica napus* [23]. It is worth noting that this rate constant also represents the average rate constant taking into account, in addition to the rate constant of the reaction of limonene with OH, those of sabinene ($k_{\text{sabinene-OH}} = 117 \times 10^{-12} \text{ cm}^3 \text{ molecule}^{-1} \text{ s}^{-1}$ (at 296K) [47]) and myrcene ($k_{\text{myrcene-OH}} = 213 \times 10^{-12} \text{ cm}^3 \text{ molecule}^{-1} \text{ s}^{-1}$ (at 294K) [48]), since these latter compounds can also be important monoterpenes emitted by rapeseed.

Finally, this list of compounds (Table 2) contains species known to be biogenic compounds: isoprene, monoterpenes and green leaf volatiles (GLV). This latter family consists of C₆ compounds emitted by almost every green plant upon herbivory or pathogen infection and known for their important role in plant defenses [49]. Several OVOCs appear in this list: formaldehyde, methanol, acetaldehyde, acetone, etc. OVOCs can be of anthropogenic or biogenic origins and can be primarily emitted by plants or generated from secondary reactions (acetaldehyde, acetic acid, etc.). Therefore, it is important to distinguish between rapeseed BVOCs emissions and background air emissions. In this context, we determined the concentration of VOCs related to the enclosed plant emissions as $\text{VOC}_{\text{rapeseed}} = \text{VOC}_{\text{chamber}} - \text{VOC}_{\text{ambient}}$ (Figure 2).

Table 2. List of selected compounds used for OH reactivity calculations with their respective $k_{i\text{-OH}}$ at 298 K and 1 atm.

Measurements	Protonated Theoretical Mass	Chemical Compound	Name	Rate Constant ($\times 10^{-12} \text{ cm}^3 \text{ molecule}^{-1} \text{ s}^{-1}$)	Ref.
PTRQi-ToFMS protonated masses	m31.018	(CH ₂ O)H+	Formaldehyde	8.5	[50]
	m33.033	(CH ₃ OH)H+	Methanol	0.9	[50]
	m45.033	(CH ₃ CHO)H+	Acetaldehyde	15	[50]
	m57.069	(C ₄ H ₈)H+	Butanol fragment	8.5	[50]
	m59.049	(C ₃ H ₆ O)H+	Acetone	0.18	[50]
	m61.029	(C ₂ H ₄ O ₂)H+	Acetic Acid	0.74	[50]
	m69.070	(C ₅ H ₈)H+	Isoprene	100	[50]

Table 2. Cont.

Measurements	Protonated Theoretical Mass	Chemical Compound	Name	Rate Constant ($\times 10^{-12} \text{ cm}^3 \text{ molecule}^{-1} \text{ s}^{-1}$)	Ref.
PTRQI-ToFMS protonated masses	m73.064	(C ₄ H ₈ O)H+	Butanone/ MEK	1.2	[50]
	m81.068	(C ₆ H ₈)H+	MTs fragment	165	[51]
	m83.084	(C ₆ H ₁₀)H+	Hexadiene/C ₆ GLV	69.1	[52]
	m105.066	(C ₈ H ₈)H+	Styrene	53	[53]
	m137.129	(C ₁₀ H ₁₆)H+	Monoterpenes (MTs)	165	[51]
		C ₂ H ₆	Ethane	0.24	[50]
		C ₂ H ₄	Ethene	7.9	[50]
		C ₃ H ₈	Propane	1.1	[50]
		C ₃ H ₆	Propene	29	[50]
		C ₄ H ₁₀	i-butane	2.35	[51]
		C ₄ H ₁₀	n-butane	2.35	[51]
		C ₂ H ₂	Acetylene	0.75	[51]
		(E)-2-C ₄ H ₈	Trans-2-butene	63.7	[34]
	GC-FID		1-C ₄ H ₈	1-butene	31.5
		1-C ₄ H ₈	i-butene	51.5	[34]
		(Z)-2-C ₄ H ₈	Cis-2-butene	56	[34]
		iso-C ₅ H ₁₂	i-pentane	3.65	[54]
		n-C ₅ H ₁₂	n-pentane	3.92	[55]
		CH ₂ =CH-CH=CH ₂	1,3-butadiene	58	[56]
		2-(E)-C ₅ H ₁₀	Trans-2-pentene	57.1	[56]
		1-C ₅ H ₁₀	1-pentene	35	[56]
		(CH ₃) ₂ CH(CH ₂) ₂ CH ₃	2-Me-pentane	5.2	[57]
		C ₆ H ₁₄	n-hexane	5.38	[55]
Other compounds		CO	Carbon monoxide	0.228	[51]
		CH ₄	Methane	0.0064	[51]
		NO	Nitrogen oxide	9.7	[51]
		NO ₂	Nitrogen dioxide	11.3	[51]

2.6. Data Availability

The CRM/PTR-MS system operated from 7–25 April 2017. However, final experimental conditions, including heating the sampling lines and sampling ambient air from a mast at ≈ 2.7 m height, commenced 10 April 2017. All other instruments were operational between the 10th and the 25th of April except for the NO_x analyzer, resulting from technical problems from 14–19 April. Since NO data is important to correct for the CRM artifact due to the recycling of OH radicals (NO + HO₂), only OH reactivity data corrected for the NO interference are presented. It should be noted that one to 2–3 days were dedicated at the beginning, the middle (12–13 April) and the end of the campaign, to perform some tests aiming to check the stability of the CRM system. In addition, enclosure measurements were interrupted for 2 days during the campaign (between the 15th and the 17th April 2017), since a fungicide (FILAN SC (concentrated suspension) consisting of 200 g/L boscalid and 200 g/L dimoxystrobin) application was planned for this period. Therefore, taking into account the CRM/PTR-MS availability with a 1h incoming air/4h outgoing air measurement sequence, as well as the availability of the other deployed instruments, only the results from the 20th to the 23rd of April, representing the later stage of the blooming period, are presented and further discussed.

3. Results and Discussion

In the first part of the result section, the variability of measured total OH reactivity in the outgoing chamber air ($m\text{ROH}_{\text{chamber}}$) as well as in the incoming ambient air ($m\text{ROH}_{\text{ambient}}$) is described, linked to some meteorological parameters. In a second part, calculated OH reactivity is determined in ambient air ($c\text{ROH}_{\text{ambient}}$) as well as from the rapeseed branch ($c\text{ROH}_{\text{rapeseed}}$), with a description of the individual contribution of biogenic and abiotic compounds to $m\text{ROH}_{\text{chamber}}$. The final part is

dedicated to discussing the possible origins of missing OH reactivity, based on the investigation of further masses measured with the PTRQI-ToFMS.

3.1. Total OH Reactivity

Figure 3b,c report the variability of measured total OH reactivity from the rapeseed dynamic chamber ($mROH_{\text{chamber}}$) with temperature, PAR and RH, from the 20th to the 23rd of April 2017. $mROH_{\text{chamber}}$ varied diurnally between the limit of detection and 47.6 s^{-1} , with an average of $12.0 \pm 6.7 \text{ s}^{-1}$. Higher values were measured during daytime and minimum values during night-time, following PAR and temperature profiles. The average day and night $mROH_{\text{chamber}}$ was 14.9 ± 7.1 and $8.1 \pm 3.1 \text{ s}^{-1}$, respectively. The night-time $mROH_{\text{chamber}}$ was in the same order of magnitude of the mean total OH reactivity measured in the incoming chamber air (mean $mROH_{\text{ambient}} = 7.5 \pm 2.4 \text{ s}^{-1}$). However, the latter did not show a clear diurnal cycle as in the chamber reactivity. Such findings were similar to the observations made by Ren et al. (2005) [14] during a field experiment conducted in an agricultural research farm, where averaged ambient OH reactivity was 6.1 s^{-1} with little diurnal variation. However, our results are lower than the mean values (20 s^{-1} at noon and 50 s^{-1} at day-break) recorded in ambient air by Lou et al. (2010) during the PRIDE-PRD2006 campaign at a rural site which was dominated by anthropogenic pollutants (CO, NO_x , etc.) at night and strongly influenced by local biogenic emissions of isoprene during the day [15]. Our results were also lower than the 25 s^{-1} OH reactivity reported by Nolscher et al. (2013) [18] from a dynamic branch cuvette enclosure system mounted on a Norway spruce and much lower than the OH reactivity recorded by Kim et al. (2011) from red oak, white pine, beech and red maple branch enclosures which ranged between tens and hundreds of s^{-1} [17]. Thus, recorded levels of OH reactivity in ambient air and in the dynamic chamber were in the lower range of total OH reactivity, compared to various forest ecosystems, where OH reactivity reached maximum levels between 30 and 100 s^{-1} [17,18,33,58–60].

From 20–23 April, the crop growth was relatively stable as shown by the stem length, the total leaf length and the fraction of open flowers evolution (see S4). $mROH_{\text{chamber}}$ showed to be clearly correlated to environmental abiotic factors rather than changes in plant physiology. As mentioned before, $mROH_{\text{chamber}}$ minima recorded during nights were close to the levels measured in ambient air, which indicates that at night, there was no or little contribution from the plant emissions to OH reactivity. This observation is confirmed by $\Delta mROH = mROH_{\text{chamber}} - mROH_{\text{ambient}}$ (Figure 3a), with relatively low values during the night, whereas, during the day, $\Delta mROH$ reached a maximum of 10 s^{-1} on average at midday (between 11h and 14h UTC). This difference appeared around 05:00 (UTC) when the sun rose ($\text{PAR} > 0 \mu\text{mol m}^{-2} \text{ s}^{-1}$), peaked around noon in parallel with maximal solar radiations and temperatures, and decreased in the late afternoon down to a minimum around 19:00 (UTC). The midday difference was the highest on the 21st of April ($\Delta mROH = 14.3 \text{ s}^{-1}$), when a higher daytime temperature and a more intense solar radiation were recorded inside the enclosure. These observations show that the contribution of plant emissions to OH reactivity were temperature- and/or light-dependent. It is worth noting that a higher temperature was recorded inside the chamber compared to ambient air (mean difference = $8.5 \text{ }^\circ\text{C}$, see S5), this should be kept in mind when analyzing $\Delta mROH$.

It is worth noting that an extra daytime peak of $mROH_{\text{chamber}}$ (Figure 3b) was recorded in the morning of the 21st of April (05:00–07:30 UTC) that did not appear on the other days and was also seen on $\Delta mROH$ (Figure 3a). This peak of $\Delta mROH$ is likely due to the subtraction of a $mROH_{\text{ambient}}$ (an average of 7.5 s^{-1}) not representative of the atmospheric conditions at that hour of the day; indeed, we remind that $mROH_{\text{ambient}}$ was only recorded 1 h each 4 h, and that the ambient air composition can change quickly. In addition, higher night-time levels were observed during the nights of 21st and the 22nd April (around 01:00 UTC). These events are further discussed in Section 3.2 while investigating the contribution of measured compounds to total measured OH reactivity.

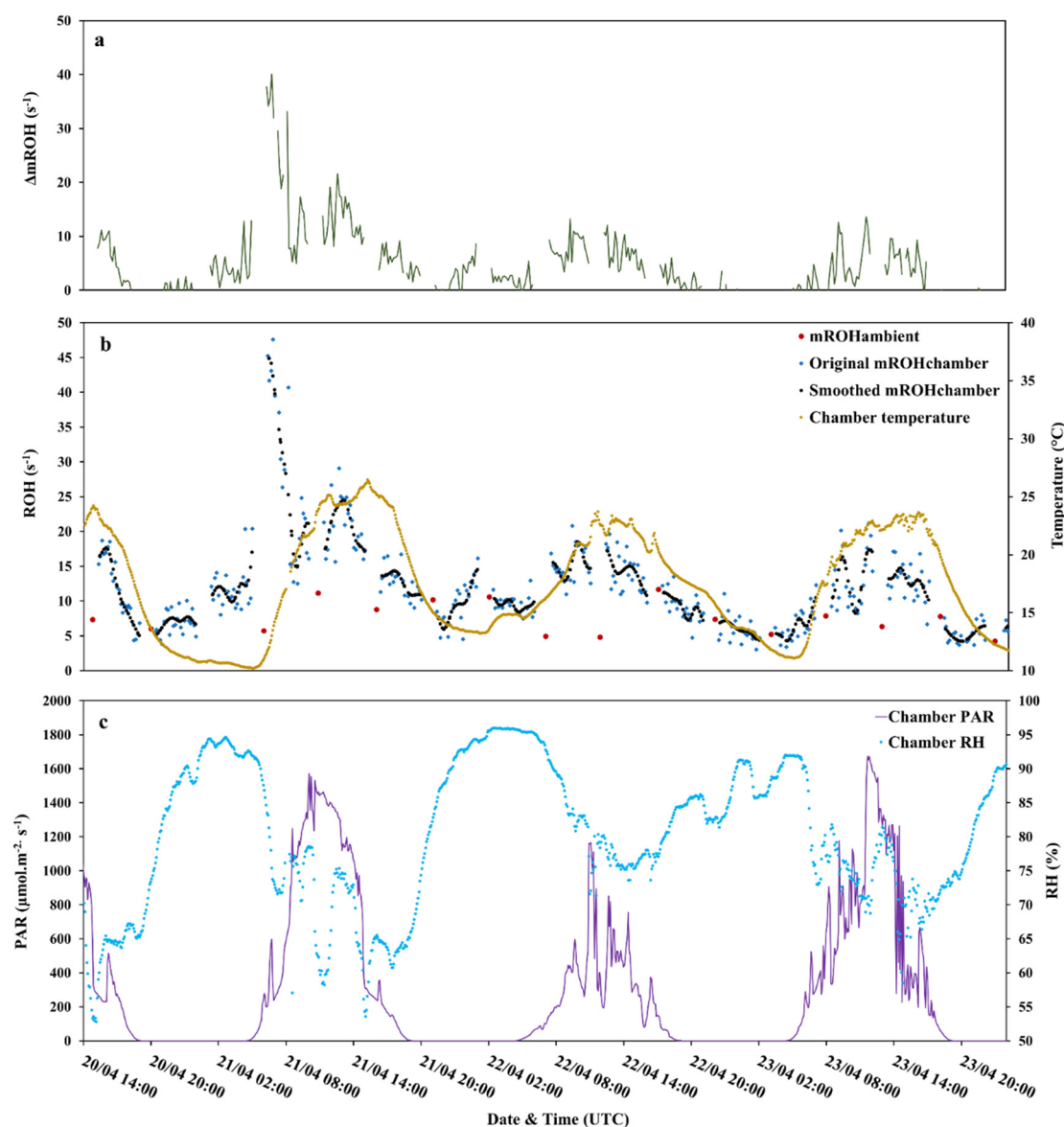


Figure 3. Variability of (a) $\Delta mROH = mROH_{chamber} - mROH_{ambient}$ and (b) total $mROH_{chamber}$ (blue dots represent the original OH reactivity data and black dots represent the trend obtained after smoothing the original data) and total $mROH_{ambient}$ (red dots) from the 20th to the 23rd April 2017 with temperature (T ($^{\circ}C$), yellow), (c) photosynthetic active radiation (PAR $\mu mol \cdot m^{-2} \cdot s^{-1}$, purple) and relative humidity (RH%, light blue), all three parameters measured inside the chamber.

3.1.1. Ambient Air Measurements and their Calculated OH Reactivity ($cROH_{ambient}$)

Measurements in ambient air included NO, NO₂, CO, CH₄, C₂–C₆ hydrocarbons and other VOCs (Figure S6a). As already stated, NO was around 0.7 ± 0.5 ppbv on average for the considered period, except for one peak recorded on the morning of the 21st April, 2017 when the NO mixing ratio reached 46 ppbv around 06:00 UTC. NO₂ showed higher mixing ratios than NO and was around 12 ppbv on average. CO was on average 151 ± 21 ppbv with a higher mixing ratio on the morning of the 21st April, occurring at the same time as the NO peak and reaching 243 ppbv around 07:00 UTC. These high peaks of NO and CO, observed on the morning of the 21st April, correspond to the hours of the day when the wind was mainly coming from the east of the site (wind direction $\approx 50^{\circ}$) and had a low speed, which suggests little mixing in the surface layer. These peaks lasted for 2–3 h and they were most likely related to traffic from the road nearby and advection from Paris and its suburbs. In terms

of calculated OH reactivity, these two compounds, accounted for 4.6 s^{-1} on average to the ambient measured OH reactivity, with a maximum of 14.3 s^{-1} on the morning of the 21st of April 2017 (S6b).

Regarding VOCs, CH_4 presented an average of 2100 ppbv, with a higher level on the night between the 20th and the 21st of April and a high/sharp peak around 01:00 UTC of the 22nd. The same observation was made on the sum of $\text{C}_2\text{-C}_6$ hydrocarbons (GC-FID) and the oxygenated $\text{VOCs}_{\text{ambient}}$ (PTRQI-ToFMS). In terms of OH reactivity, CH_4 , non-methane hydrocarbons in the $\text{C}_2\text{-C}_6$ fraction and OVOCs accounted, each, for $0.3\text{--}0.4 \text{ s}^{-1}$ on average, with a maximum of 0.7, 1 and 1.2 s^{-1} , respectively, around midnight on the 22nd. Monoterpenes accounted for 0.7 s^{-1} on average and isoprene had a negligible contribution, around 0.1 s^{-1} , both showing surprisingly the same midnight peak reaching 3.2 and 0.7 s^{-1} , respectively. This peak coincides with a wind mainly coming from the north of the site and highlights possible emissions from a nearby waste treatment plant [61]. All the mentioned compounds accounted for 6.5 s^{-1} on average of $\text{mROH}_{\text{ambient}}$, representing a relative mean contribution of 87%.

It is worth pointing out again that, assuming that the inorganic compounds and light hydrocarbons were not emitted nor deposited in the rapeseed chamber, all the compounds present in the incoming chamber air (ambient air) contribute to $\text{mROH}_{\text{chamber}}$. This contribution was around 89% and 41% during night-time and daytime, respectively.

3.1.2. BVOCs from the Rapeseed Plant ($\text{VOCs}_{\text{rapeseed}}$) and Their Calculated OH Reactivity ($\text{cROH}_{\text{rapeseed}}$)

As mentioned before, the PTRQI-ToFMS was the only instrument measuring VOCs from the dynamic chamber ($\text{VOCs}_{\text{chamber}}$) as well as in ambient air ($\text{VOCs}_{\text{ambient}}$). Thereby, it was possible to characterize VOCs originating directly from the enclosed plant ($\text{VOCs}_{\text{rapeseed}} = \text{VOCs}_{\text{chamber}} - \text{VOCs}_{\text{ambient}}$), on which we focus in the following section.

The summed mean mixing ratios of the 11 $\text{VOCs}_{\text{rapeseed}}$ is presented in Figure 4a. It exhibits a diel variability with a maximum of 11 ppbv on average around midday (11 h–14 h UTC) and a minimum during the night (\approx LOD), similar to temperature and light profiles. This observation confirms that BVOCs emissions from rapeseed and their concentrations in the enclosure are generally light- and/or temperature-dependent. It also highlights that these compounds are the main drivers of the day/night variability seen on $\text{mROH}_{\text{chamber}}$.

Between the 20th and the 23rd of April, biogenic OVOCs (listed in Table 2) accounted for the major (92%) part of $\text{VOCs}_{\text{rapeseed}}$. Among the measured biogenic OVOCs, methanol was dominant with 1.4 ppbv on average, representing 38% of the oxygenated compounds emitted, followed by acetaldehyde and acetone, with 0.8 ppbv each, accounting for 21–22%. Acetic acid (0.4 ppbv) presented around 11%, whereas the others (formaldehyde, butanol and MEK) did not exceed 8% together. Regarding isoprene and monoterpenes, they had relatively low contributions to the total $\text{VOCs}_{\text{rapeseed}}$, with 0.2 and 0.01 ppbv on average, amounting for 5.2% and 0.2%, respectively. These observations during the blooming period are in agreement with other studies on rapeseed. König et al. (1995) [62] classified rapeseed as an OVOC emitter with other crop species. Müller et al. (2002) [25] reported an important emission and/or rapid formation of carbonyl compounds from flowering rape as an important source of biogenic OVOCs in the studied ecosystem. Furthermore, these relatively high levels of OVOCs, recorded during the blooming period, were also observed later during the senescence period, starting at the end of April 2017, in the study conducted by Gonzaga Gomez et al. (2019) [27] on the same field site. However, few studies presented rapeseed as an important emitter of monoterpenes, which accounted for 60–90% of the total VOCs identified in spring oilseed rape, whereas non-terpenoid components, including OVOCs (acetaldehyde, acetone and 3-methyl-2-butanone) were found at low levels [23].

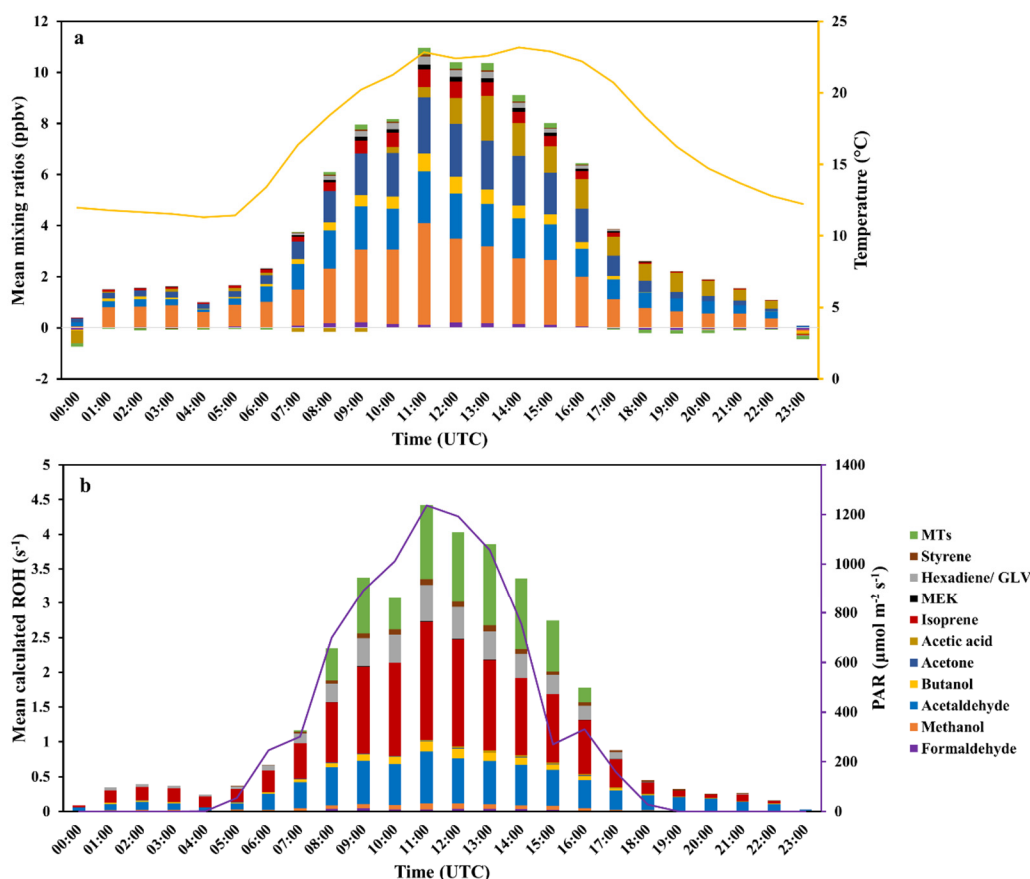


Figure 4. Averaged diurnal variation of (a) the mixing ratio of each biogenic volatile organic compound (BVOC) measured with the PTRQI-ToFMS (ppbv) and (b) their contribution to OH reactivity (s^{-1}).

When looking at individual BVOC variability (S7), all compounds had a clear diurnal cycle, except for acetic acid whose variability followed neither temperature nor light. Emissions of isoprene, as well as of the rest of the OVOCs and hexadiene/Green Leaf Volatiles (GLV), were correlated to light and/or temperature, because they exhibited higher levels during daytime compared to the night-time, and for night-time levels, they showed higher levels during the 22nd night when higher night-time temperatures were recorded. Finally, monoterpenes did not show the same behavior and their emissions were close to zero during the night, even when night-time average chamber temperature was at its highest, around 15 °C.

Figure 4b illustrates the mean contribution of each $VOC_{rapeseed}$ to the mean calculated OH reactivity ($cROH_{rapeseed}$) that exhibited an average diurnal variation between the LOD and $4.4 s^{-1}$. Interestingly, the major contributors to $cROH_{rapeseed}$ during daytime were isoprene with 40%, followed by acetaldehyde (21%) and monoterpenes (18%). Hexadiene/ GLV accounted for 12% and the rest (formaldehyde, methanol, butanol, acetone, acetic acid, MEK and styrene), did not exceed 10% all together. Despite that methanol was the most abundant compound in the chamber, its contribution to OH reactivity remains low, compared to less emitted but more reactive species, like isoprene and monoterpenes.

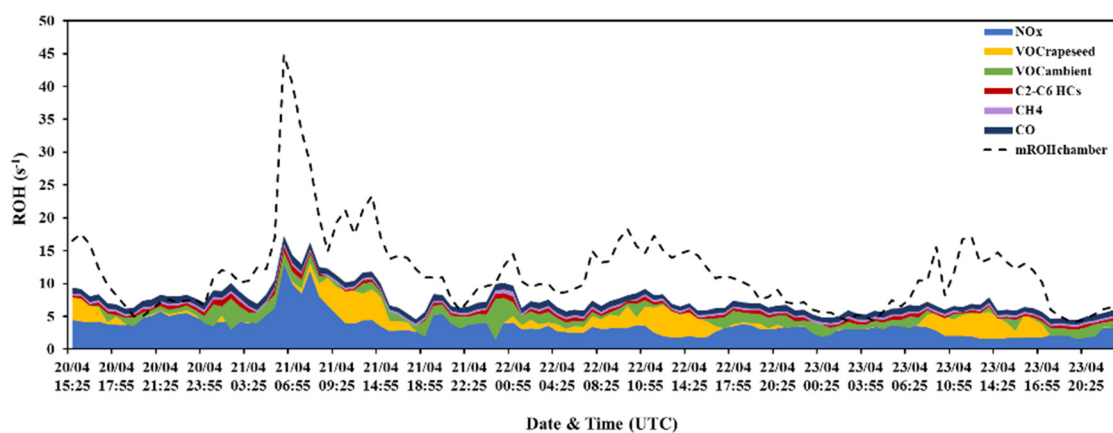
Compared to measured OH reactivity in the outgoing chamber air ($mROH_{chamber}$), all of the considered biogenic $VOC_{rapeseed}$ explained 13% on average of the $mROH_{chamber}$ during daytime with a maximum contribution at midday (11:00–14:00 UTC) reaching 24% on average, whereas during night-time, biogenic $VOC_{rapeseed}$ contribution to $mROH_{chamber}$ was negligible. Contributions of ambient species and rapeseed emissions to $mROH_{chamber}$ are summarized in Table 3.

Table 3. Summary of the mean contribution of all the measured species (same time step) in the incoming and the outgoing chamber air to $mROH_{\text{chamber}}$ and the resulting missing OH fraction.

ROH and ROH_{missing}	Daytime Avg \pm STD	Night-time Avg \pm STD
cROH NO_x (s^{-1})	3.6 ± 2.2	3.8 ± 0.9
cROH HCs C_2 - C_6 (s^{-1})	0.4 ± 0.2	0.5 ± 0.2
cROH CO (s^{-1})	0.8 ± 0.1	0.9 ± 0.1
cROH CH_4 (s^{-1})	0.3 ± 0.01	0.3 ± 0.07
cROH VOC_{ambient} (s^{-1})	0.8 ± 0.7	1.5 ± 1.0
cROH VOC_{rapeseed} (s^{-1})	1.9 ± 1.7	<LOD \pm 0.7
Mean cROH _{chamber} (s^{-1})	7.8 ± 2.6	7.0 ± 1.5
Mean $mROH_{\text{chamber}}$ (s^{-1})	14.4 ± 7.0	8.0 ± 2.6
Mean missing ROH (s^{-1})	6.6 ± 5.0	1.0 ± 1.7
Mean relative missing ROH (%)	46	13

3.2. Investigation of the Missing OH Reactivity

The comparison between measured OH reactivity ($mROH_{\text{chamber}}$) and the reactivity calculated by the sum of individually measured reactive compounds ($cROH_{\text{chamber}}$) enabled us to assess the known reactive carbon budget and to determine the unaccounted for OH sink, known as missing OH reactivity (ROH_{missing}). Figure 5 shows the variability of $mROH_{\text{chamber}}$ and $cROH_{\text{chamber}}$. Little or no missing OH reactivity was observed during the night, whereas a more important difference was seen around midday (mean midday missing OH reactivity $\approx 8 s^{-1}$), suggesting that this missing OH reactivity was mainly related to light- and/or temperature-dependent compounds either primarily emitted or secondarily formed in the chamber.

**Figure 5.** $mROH_{\text{chamber}}$ and $cROH_{\text{chamber}}$ from individual calculated ROH of VOCs (PTR-MS), VOCs (GC-FID), NO_x , CH_4 and CO summed up and the resulting missing ROH.

Among the approaches used to explore the origin of missing reactivity, one consists of correlating the missing OH reactivity with temperature (Figure 6). As such, we tried to fit the missing OH reactivity with the equation applied by Di Carlo et al. (2004) [63], adapted from the one normally used to describe temperature-dependent emissions of monoterpenes [64].

$$ROH_{\text{missing}}(T) = ROH_{\text{missing}}(293)\exp(\beta(T-293)) \quad (4)$$

with $ROH_{\text{missing}}(T)$ and $ROH_{\text{missing}}(293)$ representing the missing OH reactivity at a given temperature $T(K)$ and 293 K, respectively, and β (K^{-1}), the temperature sensitivity factor.

The β value obtained was around $0.14 K^{-1}$ with a correlation coefficient (R^2) of 0.8. In other studies, β ranged between 0.054 and $1.144 K^{-1}$ suggesting that in this study the daytime missing

OH reactivity may be related to temperature-dependent monoterpene-like emissions. Furthermore, missing OH reactivity moderately correlated with PAR, following an exponential relationship with an R^2 of 0.7, which suggests that the missing OH reactivity may also be due to light-dependent emissions, although it is difficult to disentangle radiation and temperature effects in field conditions, since both are correlated by the energy balance of the plant.

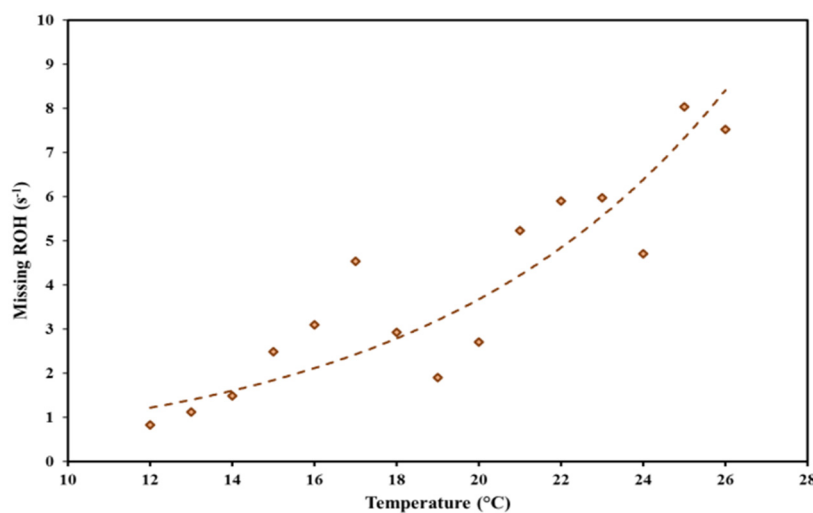


Figure 6. Variability of missing OH reactivity as function of air temperature in the chamber. The regression curve equation is $ROH_{\text{missing}} = 0.23\exp(0.14(T))$ ($R^2 = 0.8$).

Figure 5 also demonstrates the clear diurnal co-variability between $cROH_{VOC_{\text{rapeseed}}}$ (orange) and ROH_{missing} (the difference between the dashed line of $mROH_{\text{chamber}}$ and the sum of calculated OH reactivity of the individually measured species), with higher levels around midday, suggesting that primary biogenic VOC_{rapeseed} , whose emissions are driven by temperature and/or light, are potential candidates for the daytime missing OH reactivity.

Midday missing OH reactivity: In order to investigate the potential species behind the observed midday missing OH reactivity, we looked at the PTRQI-ToFMS masses that correlated up to 80% and more with isoprene. These correlations were done based on the fact that isoprene is a primary compound, which emission is known to be light- and temperature-dependent. As a result, 201 additional ions were investigated (Table S8). However, when it came to estimate their contribution to total OH reactivity, this list was reduced to only 9 masses which were tentatively identified and for which rate coefficients could be found (Table 4). Using their mixing ratios related to plant emissions (VOC_{rapeseed}), the selected 9 compounds exhibited a diurnal cycle with a maximum contribution of 1 to 1.2 s^{-1} around noon, and a negligible contribution during night hours (Figure 7). This observation is an evidence that midday missing OH reactivity can be related to unaccounted primary emitted compounds, but the identified compounds only explained a part of the missing reactivity (12% on average of midday missing OH reactivity). It is worth noting that, among the investigated ions (Table S8), m/z 41.038 was identified, as mentioned before (Section 2.5), as a potential fragment of isoprene. Considering its mixing ratio in the calculation of isoprene's OH reactivity induces a maximum increase of 3 s^{-1} and an average increase of 1.9 s^{-1} around midday, it explains 24%, on average, of midday missing OH reactivity. Thus, this test (S9) should be kept in mind while calculating OH reactivity and investigating the missing OH reactivity.

Considering the masses that correlated 90% or more with m/z 69.070 (excluding the masses already investigated and mentioned in Tables 2 and 4), their summed mixing ratios in the chamber air out ranged between the LOD and 0.8 ppbv. If all of them had a comparable reactivity to that of isoprene towards the hydroxyl radical, their contribution to total OH reactivity would reach a maximum of 2.1 s^{-1} and would explain 16% on average of midday missing OH reactivity. This contribution would

reach a maximum of 0.02 s^{-1} only if all the investigated compounds had a reactivity comparable to that of methanol, explaining only 0.1% on average of midday missing OH reactivity. Regarding the masses that correlated between 80% and 90% with m/z 69.070, their summed mixing ratios accounted for a maximum of 2.7 ppbv, contributing up to 6.9 s^{-1} and 0.06 s^{-1} , if all of them had a comparable reactivity to that of isoprene or methanol, respectively. These masses would explain up to 52% and only 0.5% on average of midday missing OH reactivity with a reactivity as high as that of isoprene or as low as that of methanol, respectively.

Table 4. List of the 9 selected masses which presented a correlation of 80% and more with isoprene (m/z 69.070).

Protonated Theoretical Mass	Chemical Formula	Chemical Compound	Rate Constant ($\times 10^{-12} \text{ cm}^3 \text{ molecule}^{-1} \text{ s}^{-1}$)	Max Mixing Ratio (ppbv)	Max Calculated ROH (s^{-1})	Reference
57.033	$\text{C}_3\text{H}_4\text{O}$	Propenal	19.6	0.23	0.11	[34]
85.064	$\text{C}_5\text{H}_8\text{O}$	Ethylvinylketone	36 ^a	0.06	0.06	[56]
85.1	C_6H_{12}	Hexene	37 ^a	0.18	0.17	[56]
87.078	$\text{C}_5\text{H}_{10}\text{O}$	2-Methylbutanal ^b	32.8 ^c	0.06	0.05	[65]
97.098	C_7H_{12}	Methylcyclohexene	96.1 ^d	0.3	0.66	[66]
109.1	C_8H_{12}	4-Ethenylcyclohexene	63.9 ^a	0.23	0.36	[56]
119.082	C_9H_{10}	Indan	19.1 ^e	0.02	0.01	[67]
133.098	$\text{C}_{10}\text{H}_{12}$	Benzene, (2-methyl-1-propenyl)-	33	0.01	0.01	[68]
157.105	$\text{C}_{12}\text{H}_{12}$	Dimethylnaphthalene	74.9 ^f	0.03	0.06	[69]

^a: Theoretical rate constant at 298 K and 1 atm; ^b: Other compounds may exist, 3-Methylbutanal and 2-pentanone; ^c: The rate coefficient of the reaction of *sec*- $\text{C}_4\text{H}_9\text{CHO}$ (2-Methylbutanal) was considered and this rate constant is higher than that of 2-pentanone ($k_{\text{OH}} = 4.56 \times 10^{-12} \text{ cm}^3 \text{ molecule}^{-1} \text{ s}^{-1}$ [70]); ^d: Rate constant at 305 K; ^e: Rate constant at 297 K; ^f: Rate constant of 1,3-dimethylnaphthalene, which is the highest among the rate constants of dimethylnaphthalene isomers with OH and which range between $57.9 \times 10^{-12} \text{ cm}^3 \text{ molecule}^{-1} \text{ s}^{-1}$ and $74.9 \times 10^{-12} \text{ cm}^3 \text{ molecule}^{-1} \text{ s}^{-1}$.

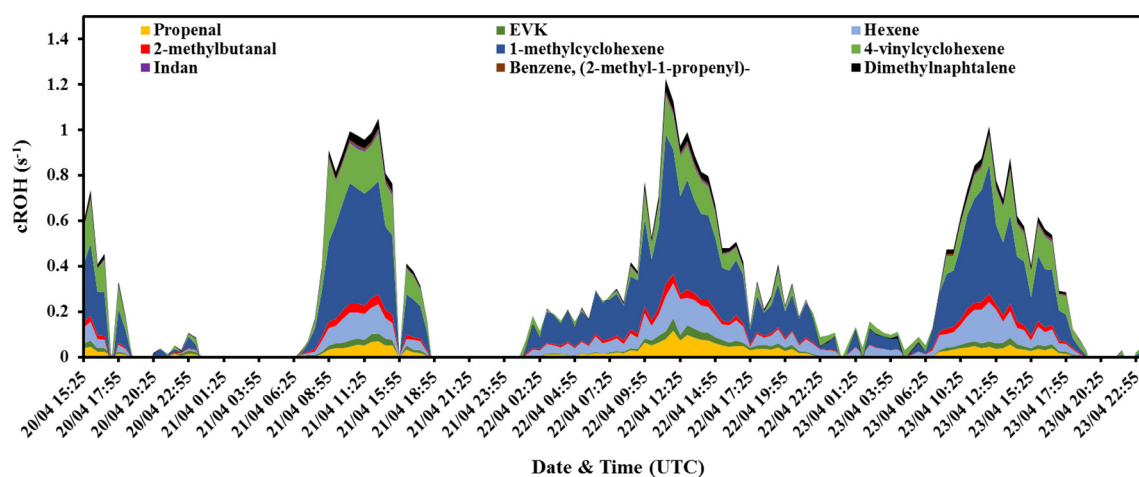


Figure 7. Summed calculated OH reactivity of the 9 additionally selected compounds. The mixing ratios used for the calculation correspond to the ones from the rapeseed branch ($\text{VOC}_{\text{rapeseed}}$) and the $k_{i\text{-OH}}$ are the ones mentioned in Table 4.

Other masses: Gonzaga Gomez et al. (2019) [27] reported the presence of other masses which were not necessarily observed in previous studies on *brassica napus* [25,62,71,72]. Among these masses, m/z 43.018 exhibited relatively large mixing ratios, reaching 5 ppbv. This observation is not surprising, knowing that this mass represents a fragment of several oxygenated compounds. This implies that the mixing ratios of some oxygenated compounds were underestimated, leading to an underestimation of the calculated OH reactivity. Based on an intercomparison exercise (Eurochamp, Orléans, 2019), in which the deployed PTRQi-ToFMS participated along with many other PTR-MS instruments, the

fragmentation rates on m/z 43.018 on the deployed PTRQI-ToFMS were the following: acetone 3.4%, acetaldehyde 3.6%, acetic acid 20%, MVK 13% and MEK 1.0% (Lafouge F., personal communication). Taking these rates into account and correcting the mixing ratios of acetone, acetaldehyde, acetic acid and MEK for potential fragmentation in OH reactivity calculations increased the calculated OH reactivity by 0.05 s^{-1} maximum. This result highlights that not accounting for fragmentation that can occur on some oxygenated VOCs is not responsible for the observed missing OH reactivity.

Other missing OH reactivity events: In addition to midday missing OH reactivity, a higher missing OH reactivity was also observed in the early morning of the 21st of April (around 07:00 UTC) and in the beginning of the evening (around 18:00 UTC) as well as at midnight and during the night of the 22nd of April 2017 (Figure 8). Regarding the night of the 22nd April, higher night-time mixing ratios of some BVOCs in the chamber were recorded (i.e., isoprene), meaning that this event could be explained by unknown biogenic emissions from the plant. Interestingly, acetic acid, whose emission did not follow clearly temperature or light, also exhibited larger concentration levels during this night. Studies showed that acetic acid can be primary emitted by vegetation or generated from secondary reactions. Gonzaga Gomez et al. (2019) [27] reported emissions of acetic acid from rapeseed during the senescence period. Furthermore, Veromann et al. (2012) [71] reported higher acetic acid emissions from rapeseed after nitrogen fertilization, which could be the case in our experiment, since N-treatment was applied twice before the campaign. Seeing the relatively short residence time and the relatively rapid air renewal in the dynamic enclosure, the chances for acetic acid formation via secondary chemistry are low. These post N-treatment emissions could also be detected under several masses (Table S8) which were identified as N-containing compounds [73]. This includes m/z 100.074, m/z 116.074, m/z 124.117, m/z 138.059 and m/z 252.158, identified as $\text{C}_5\text{H}_9\text{NO}$, $\text{C}_5\text{H}_9\text{NO}_2$, $\text{C}_8\text{H}_{13}\text{N}$, $\text{C}_7\text{H}_7\text{NO}_2$ and $\text{C}_{14}\text{H}_{21}\text{NO}_3$, respectively.

No specific behavior related to BVOCs emissions from the enclosed plant could explain the higher missing OH reactivity observed in the early morning of the 21st April (around 07:00 UTC) and in the beginning of the evening (around 18:00 UTC) as well as at midnight of the 22nd April (Figure 8). However, investigating known anthropogenic VOCs, such as benzene (m/z 79.053) and benzaldehyde (m/z 107.049), detected with the PTRQI-ToFMS, demonstrated the higher ambient mixing ratios of these compounds during the mentioned events. Therefore, these events of missing OH reactivity could be mainly attributed to emissions of anthropogenic origin.

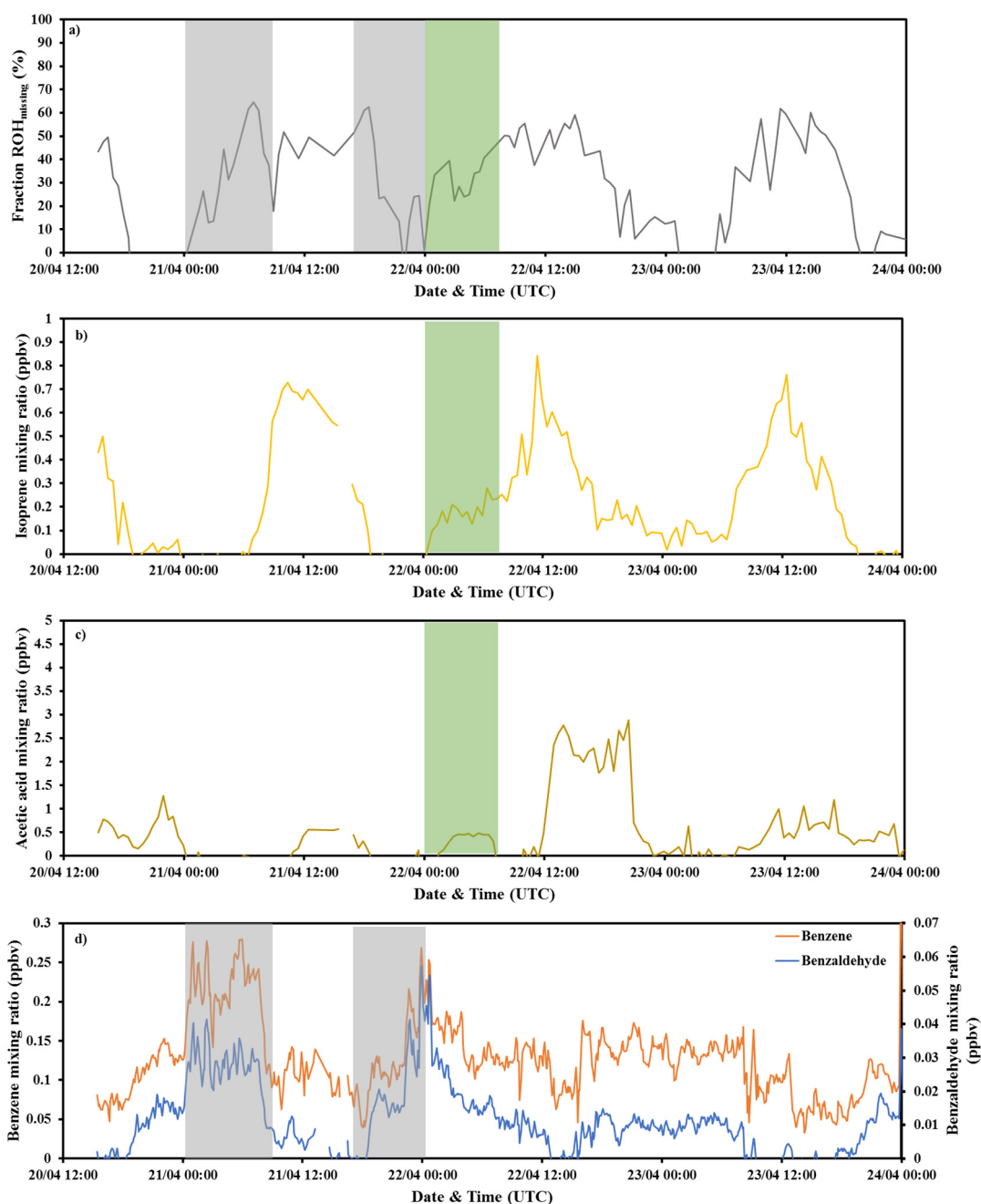


Figure 8. (a) Variability of relative missing OH reactivity ($ROH_{\text{missing}}/mROH_{\text{chamber}}$) with (b) Isoprene (rapeseed), (c) Acetic acid (rapeseed) and (d) Benzene (orange) and Benzaldehyde (blue) in ambient air. Green areas indicate missing OH reactivity mainly due to biogenic emissions and grey areas indicate missing OH mainly linked to anthropogenic emissions.

4. Conclusions

Hydroxyl radical reactivity, defined as the total loss rate of the hydroxyl radical in the atmosphere, has been shown to be an effective parameter to evaluate the loading of reactive species in ambient air. Despite the fact that total OH reactivity measurements have been performed in various environments, little has been done over agricultural ecosystems. This study presents the first characterization of total OH reactivity from rapeseed (*Brassica napus*) by means of a dynamic enclosure chamber during the blooming period in the Fr-Gri ICOS site, located in southwestern Paris during April 2017.

The average total OH reactivity measured from the dynamic chamber during the daytime was $15 \pm 7 \text{ s}^{-1}$, against $8 \pm 3 \text{ s}^{-1}$ during the night-time. The OH reactivity varied diurnally in a similar manner as BVOCs emissions, which were driven by temperature and/or light. Rapeseed emissions were dominated by oxygenated compounds, which is in agreement with some previous studies. However, in terms of OH reactivity, isoprene had the largest contribution during the day, followed by acetaldehyde and monoterpenes, whereas during the night, biogenic VOCs had a negligible contribution to OH reactivity. This provides the rationale to measure OH reactivity in addition to VOC abundance. No significant difference was observed between measured and calculated OH reactivity during night-time, whereas a missing fraction of up to 8 s^{-1} on average was observed around midday. During daytime, missing OH reactivity was exponentially correlated mainly to temperature but also to light. Further investigation of PTRQi-ToFMS masses suggested that daytime missing OH reactivity fraction is mostly related to primary emitted compounds, especially because the short residence time in the chamber (4 min) reduces the chances for secondary chemistry.

One main strength of this field experiment is to have carried out parallel measurements with the CRM and the PTRQi-ToFMS, which reflected the presence of unaccounted for reactive species. However, we were limited in terms of missing OH reactivity estimation, in particular due to challenges in identifying the chemical compounds behind the detected ions as well as their rate coefficients for the reaction with OH. This study demonstrated the need for coupling PTR-MS and GC techniques in order to identify the compounds behind the masses detected by the PTR-MS, especially when several possible compounds are suggested for the same molecular mass. Finally, it could be of great interest to perform more laboratory work aiming to broaden our knowledge on the gas-phase chemistry and kinetics of a wider variety of atmospheric compounds with the hydroxyl radical, such as for some N-containing compounds ($\text{C}_5\text{H}_9\text{NO}$, $\text{C}_5\text{H}_9\text{NO}_2$, $\text{C}_8\text{H}_{13}\text{N}$ and $\text{C}_7\text{H}_7\text{NO}_2$), which exhibited a diurnal cycle in the chamber air out.

Our work highlights that the studied rapeseed crop is a relatively low contributor to the amount of reactive species released in the atmosphere, and a relatively low contributor to total OH chemistry compared to forest ecosystems. If this is confirmed, this may mean that the expansion of bioenergy crops, like rapeseed, may have a limited impact on atmospheric OH chemistry. We should bear in mind, though, that bioenergy crop production has other impacts on the atmosphere and terrestrial ecosystems through ammonia and nitrous oxide emissions or nitrate leaching, and represent critical food security issues.

Supplementary Materials: The following are available online at <http://www.mdpi.com/2073-4433/11/3/261/s1>, Figure S1a. Linear least square fit of ΔC_2 (ppbv) vs. $\Delta(\text{m}37/\text{m}19)$ for the tests conducted on the field to assess the correction for humidity differences between C2 and C3. Green dots and red dots correspond to tests performed with ambient air and outgoing chamber air, respectively, Figure S1b. Experimental parameterization to correct for the NO interference, representing changes in C3 ($\Delta\text{C}_3 \text{ ppb} = \text{C}_{3\text{expected}} - \text{C}_{3\text{measured}}$) as function of NO mixing ratios (ppbv) in the reactor, Figure S1c. Measured ROH vs. the theoretical ROH for propane and isoprene, introduced at different concentrations and under different pyrrole/OH ratios. Circles and diamonds correspond to propane and isoprene tests, respectively, Figure S2. Schematic of the CRM/PTR-MS sampling system, Table S3. Standard Ionicon transmission curve used for computing the mixing ratio, Figure S4. Rapeseed branch evolution in the dynamic enclosure. The left vertical axis represents the length of the stem, total leaves and total flowers (in cm). The right vertical axis represents the percentage of open flowers. The red line indicates the day when the chamber was re-installed after the planned fungicide application, Figure S5. Temperature recorded inside the dynamic chamber (yellow) and in ambient air (green) during the measurement period between the 20th and the 23rd, April 2017, Figure S6.a. Variability of NO_x (NO , NO_2), CO , CH_4 , propane, propene, acetaldehyde and acetone in ambient air, with wind direction and wind speed, Figure S6.b. Variability of calculated OH reactivity of NO_x , CO , CH_4 , total HCs $\text{C}_2\text{-C}_6$ and total $\text{VOCs}_{\text{ambient}}$ (PTRQi-ToFMS), Figure S7. Variability of calculated OH reactivity of some biogenic VOCs emitted by the rapeseed plant ($\text{VOCs}_{\text{rapeseed}} = \text{VOCs}_{\text{chamber}} - \text{VOCs}_{\text{ambient}}$), Table S8. List of the 201 masses that correlate for 80 % and more with isoprene (m/z 69.070), Figure S9. Variability of calculated OH reactivity using PTR-MS data with a correction of the isoprene signal (m/z 69.070) by a factor of 1.3 (blue) and by summing the m/z 69.070 with the potential fragment on m/z 41.038 (yellow).

Author Contributions: Methodology S.B., F.T. and R.S.-E. for the CRM-PTRMS, V.G. and D.B. for the GC-FID, F.L., R.C., L.G.G. and B.L. for PTRQi-ToFMS, P.B. for the NO_x measurements and meteorological data, C.B. for the dynamic chamber, F.T. for CO measurements; formal analysis, S.B., F.T., V.G., F.L., R.C., P.B., J.K., L.G.G. and B.L., data curation, S.B., F.T., F.L. and B.L.; writing—original draft preparation, S.B.; writing—review and editing,

S.B., V.G., B.L., C.B., N.Z., J.K., R.C., P.B.; supervision, V.G.; project administration, B.L.; funding acquisition, B.L. All authors have read and agreed to the published version of the manuscript.

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Data Availability: Data are available upon request from the corresponding author.

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