Study of biogenic volatile organic compound emissions and depositions over a mixed temperate forest by PTR-TOF-MS and eddy covariance

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OBJECTIVES

Due to technical limitations, BVOC traditionally measurements were limited to a few dominant BVOC species, mostly emitted by vegetation. This study aims at providing a more detailed and complete overview of **BVOC** bidirectional exchanges over a mixed temperate forest site, even for emitted/deposited compounds less key role in which can play a atmospheric chemistry due to their high reactivity.

EXPERIMENTAL SITE AND MEASUREMENTS

- Vielsalm station, Belgium (BE-Vie): mixed temperate forest equipped with a flux tower (part of the ICOS network)
- Measurement campaign from May until October 2022
- sampling point at 51 m for $BVOC + O_3$ flux measurements and 6 additional points for BVOC + 0_3 profile measurements 4 sonic anemometers at 6, 24, 36 and 51 m above ground level Ancillary measurements (meteorological + phenological variables)



DATA PROCESSING



1 PTR-TOF-MS measurements

- PTR-TOF-4000, Ionicon Analytik GmbH
- 10 Hz measurements with E/N = 136 Td
- Background meas. every 4 hours
- Calibration meas. ± every 5 days

2 Peak detection & peak area quantification

- Carried out with IDA (Ionicon Data Analyser) software
- 3 m/z peaks clustering
- m/z values can slightly shift between IDA runs \rightarrow need to unify the detected m/z correlated to the same signal
- **Density Based Clustering in Application** with Noise (DBSCAN) algorithm to identify m/z clusters
- Optimization of the trade-off between fraction of m/z values not assigned to a

4 Fluxes computation: critical steps

- Tool based on InnFlux (Striednig et al., 2020), transcripted in Python
- Lag time determined for isoprene (m/z 69.069) and applied to other m/z
- Spectral correction factors determined from the comparison of sensible heat and isoprene cospectra, then applied to other m/z. Yields a half-power cut-off
- **5** Detection of significant exchanges **6** Compound attribution
 - Methodology:
- A. Consider half-hour (HH) significant if: |flux| > |LOD|
- B. Consider the day significant if the ratio of significant HH to the number of available HH during daytime (8 to 20h) is superior to 0.125 (i.e. at least 3 of the 24 HH during daytime are significant).
- Analysis of the fluxes intercorrelations to detect pure fragments, (water) clusters and isotopes
- Attribution of a chemical formula and a compound name to parent ions
 - Based on Pagonis et al. (2019) and Yáñez-Serrano et al. (2021, GLOVOCS) databases

- Runs on mass spectra acquired in the course of a day
- Provides the detected mass to charge ratios (m/z) and their peak areas (in normalized counts per second, ncps)
- Up to 820 m/z detected per day

cluster, and fraction of clusters containing multiple m/z peaks for a single IDA analysis

- **Clusters considered for further analysis** based on constraints on the cluster width and the fraction of data above the limit of detection for 20s aggregates
- 224 m/z selected after clustering

frequency of 0.14 Hz and a flux correction factor of about 1.10 for the mean wind speed.

- Flux limits of detection (LODs) computed at the 99% confidence level: $LOD = 3 \cdot random \ error$
- Stationarity tests not considered relevant for this dataset
- Consider the m/z value significantly 7 Flux calibration exchanged if it shows at least 3 consecutive significant days over the campaign (consistent exchange).
- Allows for the detection of episodes of short exchanges
- This methodology detects 69 m/z with significant exchanges consistent in time

For compounds not contained inside the calibration bottle, calibration factors either found in Koss et al. (2018) or taken from previous measurements with a PTR-Quad-MS in similar working conditions

RESULTS

An 'ideal period' (in terms of measurement and meteorological conditions) was chosen between 2022-07-15 and 2022-08-01 for results presentation.

Figure 1 shows that all species are on average emitted for that period, except from formaldehyde and m/z 49.027 (uncertain compound attribution so far). The BVOC budget is dominated by monoterpenes, isoprene and methanol with respective percentages of the net flux for that period of 51%, 28% and 7%.

Figure 2 illustrates the clear dependence of BVOC emissions on air temperature. Moreover, methanol and ethanol show some deposition as relative humidity rises, probably linked to adsorption/desorption of these compounds in water films.



Figure 1. Bar plot of mean emissions and depositions sorted by decreasing mean emissions (in log scale) between 2022-07-15 and 2022-08-01 for calibrated compounds. Compounds with uncertain attribution are labelled as 'organic compound' in dark red.



Figure 2. Times series of 2h-mean measured variables between 2022-07-15 and 2022-08-01. (a) Air temperature and relative humidity. (b) Two compounds with highest emissions. (c) Two compounds with clear bidirectional exchanges.

PERSPECTIVES

- This poster presents the preliminary results of the 2022 measurement campaign. With the extensive dataset acquired, BVOC flux dynamics will be studied as well as their relationship with O_3 fluxes, meteorological and phenological variables.
- This BVOC + O_3 concentration and flux dataset will be used to test and possibly improve existing mechanistic models simulating the vertical surface-atmosphere exchanges of trace gases. Measures of NO_{x} concentrations should also be carried out to complete this approach.
- The acquired BVOC and sonic anemometer profiles will be used in a Lagrangian inverse modelling approach to infer sources and sinks of BVOCs within the soil-plant continuum. Such information will increase our understanding of the mechanisms controlling BVOC exchanges.

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