



Organic aerosols mixing across the tropopause and its implication for anthropogenic pollution of the UTLS

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Abstract. Increasing anthropogenic emissions have led to numerous organic compounds in the atmosphere, with uncertain effects on climate, ecosystems, and human health. Particularly, the composition and impact of organic aerosol in the upper troposphere and lower stratosphere (UTLS) remain poorly understood, with few studies addressing the general distribution of aerosols in this layer. In this work, we present a comprehensive analysis of a tropopause fold and convective systems during an airborne campaign over central Europe in summer 2024. We collected filter samples with a multichannel sampler for organic aerosol, effectively separating tropospheric and stratospheric air masses. As a result, we analyzed the chemical composition on a molecular level using ultra-high-performance liquid chromatography coupled with high-resolution Orbitrap mass spectrometry. A subsequent non-target analysis provides novel insights into compositional differences throughout the UTLS. Our findings reveal numerous anthropogenic organic compounds, including $C_8H_{19}O_5PS_2$ and $C_{15}H_{15}NO_3S_2$, alongside dicarboxylic acids, organosulfates, and oxidation products of volatile organic compounds found in stratospheric samples. Additional target analysis identifies pollutants like perfluorooctanoic acid and Tris(2-chloropropyl) phosphate that redistribute from the ground. These findings underscore the importance of transport processes to high altitudes and the growing impact of anthropogenic pollution, contributing to a better understanding of the relationship between emissions, the chemical composition of the UTLS, and climate effects.

1 Introduction

The chemical composition of the atmosphere changes as a consequence of urbanization, developing industry and anthropogenic activities (Riese et al., 2005). With regard to a significant increase of emissions (Yu et al., 2016; Lamarque et al., 2010), models suggest a growing urgency to understand its impact on climate and climate model projections (Solomon et al., 2011). As a part of the atmospheric composition, aerosols play an important role in the global atmospheric radiation budget due to their



20 influence on the radiative forcing by scattering, absorption, as well as aerosol-cloud interactions (IPCC, 2023). Especially the upper troposphere and lower stratosphere (UTLS), located between 8 and 20 km altitude in the mid-latitudes, shows a high sensitivity to changes in atmospheric composition and mixing, yet transport and feedback processes in this region remain poorly understood (Riese et al., 2012; Murphy et al., 2021).

Throughout earlier research the composition of stratospheric aerosol was thought to be mainly particulate sulfate, mostly associated with volcanic eruptions or derived from precursor species such as carbonyl sulfide and sulfur dioxide (Myhre et al., 2004; Neely et al., 2013; Weisenstein et al., 1997). However, recent studies highlight the importance of the organic fraction of UTLS aerosols, especially in the context of an increasing number of wildfires as an important source (Yu et al., 2016; Martins-son et al., 2019; Andreae et al., 2018; Benoit et al., 2023; Dumelié et al., 2024).

Organic matter makes up to 50 % of stratospheric aerosol (Murphy et al., 2013; Froyd et al., 2009) and is either being formed by gas-to-particle conversion (Hallquist et al., 2009) or transported to high altitudes from lower tropospheric sources (Martinson et al., 2019). Organic aerosols hereby can be directly emitted as primary aerosols or originate from the atmospheric oxidation of volatile organic compounds (VOC) leading to secondary organic aerosol (SOA) (Goldstein and Galbally, 2007). In general aerosols can be of biogenic or anthropogenic origin, where thousands of compounds contribute to a variety of reactions, resulting in different chemical composition, sizes and therefore various properties of aerosol particles (Lippmann, 2011). In particular, organic aerosol present within the UTLS can be transformed by oxidation, cloud-processing and deposition, although detailed processes are still unknown (Jimenez et al., 2009; Pöschl, 2005).

In-depth studies of organic aerosols in the UTLS by online measurements or offline high-resolution mass spectrometry are limited so far, despite a few novel observations (e.g. Benoit et al. (2023)). So far, trace gases, humidity and non-refractory submicron aerosol composition were measured within the UTLS (Köppe et al., 2009; Joppe et al., 2024; Schmale et al., 2010). Since the organic fraction of UTLS aerosol is highly complex and not well understood, source apportionment, transport mechanisms, effects on the radiative forcing as well as the anthropogenic influence are insufficiently represented albeit preliminary studies hint at their importance for modelling current and future climate states. For this reason, a comprehensive understanding on the molecular level is needed to accurately assess the anthropogenic impact, understand microphysical and chemical properties and improve climate predictions as well as facilitate the development of more targeted studies.

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In this study, we target the question which differences in chemical compositions of aerosols occur throughout the different regions of the UTLS. Especially the differences above and below the tropopause are of major interest, to further understand transport and mixing. In the following, we present two case studies from the tropopause composition gradients and mixing experiment (TPEx) campaign in June 2024, probing the different regions of the UTLS (Bozem, 2025). In comparison to that, we implemented one ground measurement stationed at the rural site Kleiner Feldberg, to contrast the ground based with free tropospheric data. With our in-house built sampler for organic aerosol particles (SOAP), we were able to collect up to five filter samples throughout one flight, including one flight blank. As a consequence of that, different regions within the UTLS can be investigated separately. We further analyzed the organic fraction, using ultra-high performance liquid chromatography (UHPLC), coupled with high-resolution Orbitrap-mass spectrometry (HRMS). This setup facilitates a non-target analysis that



allows for the identification of known as well as yet unknown organic compounds. This approach enables us to distinguish stratospheric and tropospheric compounds and detect transport between both layers. Our findings contribute to a more comprehensive understanding of the UTLS aerosol composition, potentially highlighting how anthropogenic emissions are transported across the tropopause and might affect this sensitive region of the atmosphere.

2 Methods

2.1 Measurement Flights

The TPEx campaign, which was an airborne field campaign of the TPChange collaborative research center, was conducted with the aircraft Learjet 35A. Eight scientific flights and one test flight were carried out between 03 June and 21 June 2024 over Germany, the Baltic Sea and parts of Sweden (Fig. 1). All flights took off in Hohn (54.3153° N, 9.5256° E, Schleswig-Holstein, Germany) accomplished by the Gesellschaft für Flugziieldarstellung (GFD). The collected filter samples cover an altitude range of 0.9 km to 12.1 km, with a sampling duration ranging from 15 to 140 minutes. The latitude range was 49.9° N to 58.8° N and a longitude range from 6.5° E to 14.3° E. SOAP performed on every flight, collecting three to four samples and at least one flight blank per flight. The details on the filter collection are summarized in Table S1.

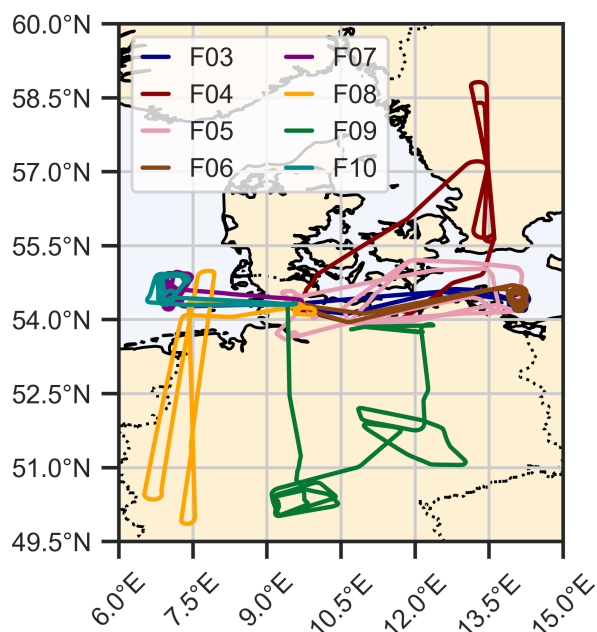


Figure 1. Overview of all flight routes during the TPEx campaign. The different flights are marked in one color each and took place in summer 2024. The exact dates are - F03: 11 June, F04: 12 June, **F05: 13 June**, F06: 14 June, F07: 17 June forenoon, F08: 17 June afternoon, **F09: 18 June**, F10: 20 June.



2.2 Data and sample acquisition

2.2.1 Sample Collection on Ground

70 Additionally to the scientific flights, we collected ambient air filter samples at the Taunus Observatory, Kleiner Feldberg (50.2219° N, 8.4468° E). The samples were collected with a high-volume sampler (DHA-80, Digitel Switzerland) on pre-heated glass fiber filters (150 mm diameter, M 160, AHLSTROM MUNKSJÖ). The sampling time was 12 h at a flow rate of 500 L min⁻¹. The particle cut off was 2.5 μm. All filter samples were sealed and stored at -20 °C after sampling.

2.2.2 Sample collection with SOAP

75 For our study, we built a multichannel aerosol sampler, which has been airborne certified for the integration in the underwing pod of the research aircraft Learjet 35A (Fig. S1). It consists of two scroll pumps that were able to draw at standard conditions in total 60 L min⁻¹ onto a quartz microfiber filter (47 mm diameter, Whatmann QM-A, cytiva). With a system of five solenoid valves which were normally closed and a bypass, we were able to collect five filter samples individually for each flight. Based on calculations according to von der Weiden et al. (2009), we estimated the particle losses, caused by the design of the inlet
80 system, to be less than 10 % in the size ranges from 10 nm to 2000 nm (Fig. S2 and S3). SOAP therefore captures particles in the Aitken and accumulation mode (Seinfeld and Pandis, 2016). Depending on the scientific aim, we sampled three to four samples, with the remaining one to two filters used as a flight blank. After each flight, filters were stored below 0 °C to prevent artefact formation (Resch et al., 2023).

2.2.3 Trace gas and aerosol properties data acquisition

85 In total 14 instruments were deployed either on board, outside the aircraft in the underwing pod or in the tropopause composition towed sensor shuttle (TOSS). The trace gases carbon monoxide (CO) and dinitrogen oxide (N₂O) were measured by university of Mainz airborne quantum cascade lasers spectrometer (UMAQS), as well as ozone (O₃) measured by the 2BTech ozone monitor. The mass concentrations of different non-refractory aerosol particles between 80 and 800 nm were acquired by the civil aircraft for the regular investigation of the atmosphere based on an instrument container-aerosol mass spectrometer (CARIBIC-AMS) with the optical particle counter (Grimm/Durag 11-S, Sky-OPC) (Schneider et al., 2025). The aerosol
90 number-size distribution was determined using an ultra-high sensitivity aerosol spectrometer (UHSAS). An overview of the deployed instruments is described by Bozem (2025).

2.3 Offline filter analysis

2.3.1 Sample extraction

95 To detect a broad range of compounds, we extracted the filters by using two different methods. A) We took two 6 mm punches and wetted them with 130 μL of 9:1 ultrapure water (MilliQ Reference A+, Merck KGaA) and methanol (0.1 micron filtered, UHPLC-MS, Thermo Fisher Scientific Inc.). Subsequently, the vials were placed on an orbital shaker (KS-15, Edmund Bühler



GmbH) at 300 rpm for 20 min. Afterwards, the solvent was extracted and filtered using a disposable polytetrafluoroethylene (PTFE) filter (pore size: 0.2 μm , macherey nagel).

100 B) We cut the remaining filter into pieces and wetted them with 2 mL methanol. After placing the vials on the orbital shaker (20 min, 300 rpm), the solvent was transferred into a sharp bottom vial. The remaining filter pieces were wetted again with 500 μL methanol and placed on the orbital shaker once more. The solvent was added into the sharp bottom vial and evaporated at room temperature using a continuous nitrogen flow. After evaporation, the remaining droplet was resolved in 80 μL 9:1 MilliQ:methanol and filtered using a PTFE filter.

105 2.3.2 UHPLC-HRMS measurements

The extracts and authentic standards were measured using an UHPLC (Vanquish Flex) combined with HRMS (Q Exactive Focus Hybrid-Quadrupole Orbitrap, both Thermo Fisher Scientific Inc.). The measurements in both positive and negative ionization mode were carried out using a heated electro spray ionization source (HESI). A C_{18} -Column (CORTECSTM T3, 2.7 μm x 150 mm, Waters) at 40 °C was used for the chromatographic separation. The separation was carried out by a gradient elution with MilliQ and methanol, both, with 0.1 % formic acid. The injection volume was set to 15 μL . More details can be found in Table S2.

As authentic standards we used: phthalic acid (99.5 %, Merck), nitrophenol (100 %, Merck SIAL), suberic acid (99 %, Fisher Scientific), azelaic acid (+98.5 %, Merck Supelco), adipic acid (99.5 %, Merck Fluka) and Tri-isobutylphosphate (99 %, Sigma Aldrich).

115 2.3.3 Non-target analysis and elemental composition group assignment

The raw data was processed using Compound Discoverer 3.3 and FreeStyle 1.8 SP2 (both Thermo Fisher Scientific Inc.) to run the non-target analysis. For this analysis, Compound Discoverer uses the retention time, mass-to-charge ratio (m/z) and isotopic patterns as well as fragmentation patterns to assign molecular formulae. Additionally, the spectra are compared with the databases mzCloud (HighChem LLC, 2013-2021) and a self-built *Aerosolomics* database in mzVault 2.3 (Thermo Fisher Scientific Inc.) by Thoma et al. (2022), to further identify secondary organic aerosol compounds. The *Aerosolomics* database contains m/z , retention time and MS^2 spectra from laboratory experiments, simulating the atmospheric oxidation of volatile organic compounds.

Apart from comparison with databases, a potential molecular formula is assigned to compounds with a sample-to-blank ratio higher than five, which refers to the flight sample and the flight blank. The elements allowed for the formula prediction are bromine (Br), chlorine (Cl), carbon (C), oxygen (O), hydrogen (H), sulfur (S), nitrogen (N) and phosphorus (P). The formulae were then classified to the elemental composition groups CHO, CHNO, CHNOS, CHOS, CHOP, CHN and *other*, if none of the previous groups applies. Compounds that could not be assigned to a sum formula are excluded from our analysis. By additionally calculating the aromaticity equivalent according to Yassine et al. (2014), the CHO group can be distinguished into CHO_a (aromatic organics) and CHO_n (aliphatic organics). The details on the workflow applied can be found in the SI S4.2.



130 All signals are blank corrected and normalized according to sampling time and flow. Finally, to declare a degree of certainty for our compound identification, we used the confidence levels established by Schymanski et al. (2014).

2.4 Trajectories

For our analysis of the air mass origin and possible transport pathways, we use kinematic trajectories calculated with the Lagrangian analysis tool LAGRANTO (Sprenger and Wernli, 2015). Every minute, 60 trajectories are started from the longitude, latitude and pressure position of the aircraft. This results in a total of 15,316 trajectories for each of the analyzed flights F05 and F09. The position of these air parcels is calculated backward in time for up to ten days. The trajectory calculation uses 3D grid-scale wind data from the operational global ICON model (Zängl et al., 2014) run at the German weather service (DWD). Analysis data for 00 UTC, 06 UTC, 12 UTC and 15 UTC is combined with forecast data for lead-times of 1–5 h to achieve an hourly resolution of the wind field data. The ICON wind field data has been interpolated from its original grid (R3B07, effective grid spacing 13.5 km) onto a regular grid with a resolution of $0.15^\circ \times 0.15^\circ$ prior to trajectory calculation. In the vertical ICON uses a SLEVE coordinate system with 120 levels with a constant spacing of 300 m between about 4 and 15 km agl.

3 Results and discussion

3.1 Flight selection for investigating the chemical composition of the UTLS

To understand the chemical composition of organic aerosol as well as possible transport processes in the UTLS, eight research flights were conducted with SOAP, sampling filters for offline analysis. Out of those we selected flights F05 and F09 as case studies, that contrast each other with respect to meteorological conditions, altitude ranges and geographical regions, to investigate and compare the chemical composition within different parts of the UTLS.

F05 took place on 13 June 2024 and represents a racetrack pattern over the Baltic Sea, with downward mixing in the first half and low Richardson numbers and a shear layer in the second half. The low Richardson numbers indicate mixing potential at the tropopause (Woods, 1969) and at a tropopause fold in the measurement region, which can be associated with a strong exchange across the usual tropopause barrier (Holton et al., 1995; Shapiro, 1980). Due to the stepwise increase in height, which covered a range from 6,000 to 10,000 m, we were able to sample the different regions for at least 40 min each.

Flight F09 was conducted on 18 June 2024 and contains a low altitude pass over Spielberg, Germany (50.3207° N, 9.2638° E), and flight tracks across Germany to capture convective systems. The low altitude flight at around 900 m allows for a direct comparison to ground measurements station at the Taunus Observatory, which is within a 60 km proximity from the ground pass. Throughout the flight, the filters were sampled in four different conditions: before the ground pass, during the ground pass, throughout convective systems and in the lower stratosphere at 12,000 m altitude.



3.2 Characterizing the UTLS region

To understand which air mass was sampled during the flight, the troposphere and stratosphere needs to be defined, which requires the determination of the tropopause e.g. from gridded analysis data. Here, the dynamical tropopause, defined by a threshold value of potential vorticity (Danielsen, 1968; Shapiro, 1978), the thermal tropopause, defined by the world meteorological organization (WMO, 1957), and the chemical tropopause, based on trace gases, can be used (Joppe et al., 2024; Müller et al., 2015; Fischer et al., 2000).

Since we focus on investigating the chemical composition and the occurrence of different compounds in different regimes, we use threshold values of different trace gases as a tool to distinguish stratospheric from tropospheric air masses. This approach is based on trace gases, which have either sources in the troposphere and are rather constant in the stratosphere or the other way around. N_2O can be used as a tracer for stratospheric air, since it has no sinks in the troposphere, a long life time and no sources in the stratosphere. Therefore, the mixing ratio is constant in the troposphere, showing a background level of 336–338 ppbv and decreases in the stratosphere. O_3 in the contrary is photochemically produced in the stratosphere, with a lifetime of up to six years, and shows relatively low abundance in the troposphere. For this reason, threshold values of 70–100 ppbv have been typically used to determine the lower boundary of the stratosphere, though local chemistry, affecting the O_3 mixing ratios can complicate its interpretation (Bethan et al., 1996; Zahn et al., 2000). CO in addition is used to determine mixing processes and evaluate tropospheric influence (Hoor et al., 2002; Fischer et al., 2000). In contrast to N_2O , CO is considered a tropospheric tracer, which concentration decreases in the stratosphere in the absence of mixing with tropospheric air. The main production of CO in the stratosphere is the temperature-dependent methane oxidation by hydroxyl radicals (OH), which is less efficient than the CO reaction with OH , serving as a sink. Because of a life time of two to three month, elevated CO concentrations indicate recent tropospheric influence on the lowermost stratosphere, since photochemical in situ production of CO is too slow to account for the elevated CO in this region (Pan et al., 2004; Hoor et al., 2002).

In this study, stratospheric air masses are defined by conservative thresholds for mixing ratios of $\text{O}_3 > 100$ ppbv and $\text{N}_2\text{O} < 330$ ppbv, to avoid interference with tropospheric air masses. In general, a clear anticorrelation of N_2O and O_3 is measured, which aligns with the expected tracer-tracer correlations. The CO concentrations show no extraordinary peaks in the stratospheric regimes, which indicates that no recent pollution or exchange events occurred. The mixing ratios measured during the flights are displayed in Fig. 2. The filter sampling periods are displayed in colored bars, and the period where stratospheric air was sampled is marked by an orange background. The sampling conditions for each filter as well as the altitude are given in Table 1. Notably, the tropopause fold captured at a constant flight level during F05 is visible by measuring tropospheric air in between stratospheric air.

3.3 Chemical composition of organic aerosols in the different regions of the atmosphere

To analyze the different chemical composition of organic aerosol throughout the UTLS region, each filter, taken at a different part of the atmosphere, was analyzed individually. The chemical composition is derived by UHPLC-HRMS and thus attributing molecular formulae to each compound, based on their exact mass. To get an overall molecular fingerprint of each sample, the



Table 1. Filter number of flight F05 and F09 with respective sampling condition and name it will be further referred to.

Filter number	Sampling condition, Altitude	Filter number	Sampling condition, Altitude
F05 - The Fold		F09 - Convection Hunting	
Filter 1	tropospheric, Troposphere 6–7 km	Filter 1	tropospheric, Troposphere 10 km
Filter 2	mostly tropospheric, Troposphere 7–10 km	Filter 2	tropospheric, Ground Pass 0.9 km
Filter 3	stratospheric, Stratosphere 10 km	Filter 3	tropospheric, Troposphere 8–12 km
Filter 4	tropospheric, Troposphere 10–8 km	Filter 4	stratospheric, Stratosphere 12 km

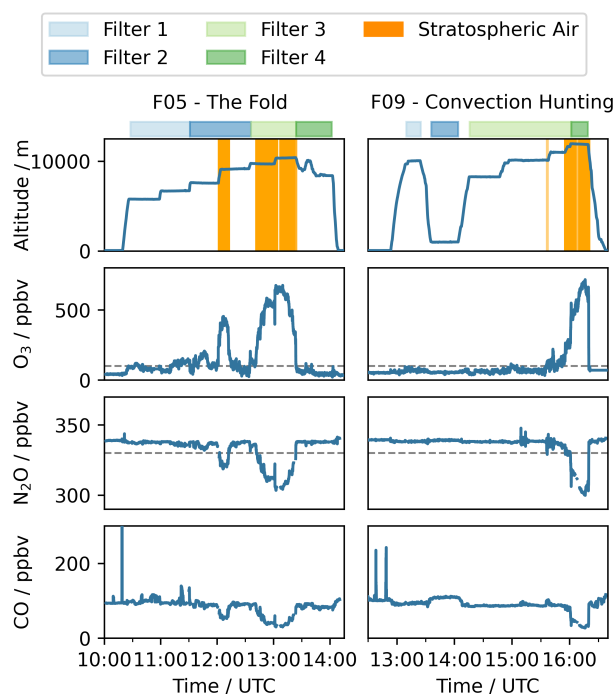


Figure 2. Overview of the altitude, measured trace gases CO, N₂O and O₃ in ppbv for flight F05 - The Fold and F09 - Convection Hunting at the respective time (UTC). The filter sampling times are displayed with colored bars with light blue: filter 1, dark blue: filter 2, light green: filter 3, and dark green: filter 4. The regimes when stratospheric air was sampled, according to tracer gas observations, are marked in orange. The condition for stratospheric air is N₂O < 330 ppbv and O₃ > 100 ppbv, marked by grey dashed lines.

compounds are assigned to elemental composition groups (CHOn, CHOa, CHNO, CHOS, CHNOS, CHOP, CHN and *other*). Figure 3 shows the relative abundance of each compound group for the respective region (exact numbers can be found in Table 2). Additionally, AMS measurements of submicron sulfate (SO₄²⁻) as well as the size distribution measured by UHSAS are compared to our observations (Fig. S4–S6).



195 For F09, the ground measurement at the Taunus Observatory from that day is also displayed, to enable a comparison with the
air masses sampled during the ground pass. It should be noted that we are mainly discussing the negative ionization mode
and that ionization as well as transmission efficiencies vary for each compound. Therefore, the signal intensities or different
groups do not directly represent concentrations of the respective compound. Still, the occurrence of the compound groups can
be compared relatively between the samples, indicating changes in the composition of organic aerosol. In general, each region
200 and each flight shows its own unique chemical composition as it will be discussed in the following sections.

CHOa and CHOn in the UTLS

Across all samples CHO has the highest contribution, splitting up into the aliphatic CHO (CHOn) and a small fraction of aro-
matic CHO (CHOa) compounds. The measurements at the ground station and the ground pass during F09 show the highest and
205 almost identical abundances, indicating a ground-based source for CHOn and CHOa compounds and served as independent
validation of the aircraft sampling. Precursors, such as volatile and semi-volatile alkanes are known to dominate biogenic and
anthropogenic emissions, followed by aromatics and unidentified oxygenated compounds (Ziemann and Atkinson, 2012). Fur-
ther up in the troposphere and with increasing altitude, their abundance decreases and therefore suggests dilution throughout
the uplifting process. This theory is supported by other aircraft based observations, described by Johansson et al. (2020) and
210 Höpfner et al. (2019). The dilution is not linear with increasing altitude, which could be due to temperature decrease and thus
condensation of intermediate volatile organic compounds. This volatility class is known to be an important source of SOA
mass in the atmosphere (Weitkamp et al., 2007; Tkacik et al., 2012; Topping et al., 2013).

CHNO, troposphere vs. stratosphere

215 When comparing the stratospheric to tropospheric measurements, the high CHNO abundance in F05 in the stratosphere stands
out prominently. As a product of VOC oxidation in presence of NO_x, which can originate from anthropogenic sources or
lightning activity, we identified some of these compounds as organonitrates (Lambe et al., 2017; Curtius et al., 2024). All
tropospheric measurements show about ten times lower CHNO abundances. Notably, only a few compounds show large signal
intensities that lead to a high CHNO abundance and will be further described in detail below. The stratospheric samples of F09
220 show in contrast to F05 a whole different chemical composition, where CHNO is not as pronounced, whereas other compounds
like CHOS, CHOP and *other* are mainly contributing to the aerosol composition.

CHOS, troposphere vs. stratosphere

The majority of CHOS are organosulfates, which are especially pronounced in the stratospheric sample of F09, as they are less
225 abundant in the respective tropospheric samples or the ground measurement. Since the formation of organosulfates is often
linked to the occurrence of sulfate (Surratt et al., 2006; Lin et al., 2011; Minerath and Elrod, 2009), the observations were
compared to the respective AMS measurements. The findings are coherent with our hypothesis, displaying an elevated sulfate
concentration of up to 0.6 $\mu\text{g m}^{-3}$ during the respective sampling time for the stratospheric sample (Fig. S4), compared to
on average 0.15 $\mu\text{g m}^{-3}$. An elevated sulfate abundance is generally expected according to earlier research by Crutzen (1976)



230 and Junge et al. (1961), observing a stratospheric sulfate layer. Moreover, previous work studying the UTLS region shows high sulfate concentrations with increasing altitude (Appel et al., 2022; Duncan et al., 2007; Tomsche et al., 2022). The same relative abundance of organosulfates is observed in F05's higher tropospheric samples. An explanation might be the influence of the tropopause fold, and the potential cross tropopause mixing at the edges of the fold (Sprenger et al., 2003). The elevated organosulfate abundance in F05 is coherent with elevated sulfate concentrations detected by the AMS (Fig. S5) throughout
235 almost the entire flight. The AMS hence reflects both observations that we made with our filter samples.

CHOP and other, troposphere vs. stratosphere

Another novel finding is the abundance of the compound group *other*, which does not seem to have any trend or altitude dependency. The ground pass during F09 and the ground station show the lowest abundances for compounds assigned to *other* and
240 F05's stratospheric sample contains almost no compounds in that compound group. For F09, one reason could be atmospheric processing like the addition of functional groups, oxidation processes or enhanced condensation due to lower temperatures that result in a decrease of volatility and thus enhancing the compounds fraction in the total organic aerosol (Atkinson and Arey, 2003; Pankow and Asher, 2008). A similar pattern is visible for CHOP, which is lowest for the ground pass and the ground station, as well as the stratospheric sample in F05. CHOP as a compound class encompasses organophosphates, which
245 are anthropogenic high-production volume chemicals, such as flame retardants. These compounds appear in the atmosphere as semi-volatile organic compounds that undergo gas-to-particle partitioning and atmospheric processing (Liu et al., 2021). Compounds assigned to *other* are e.g. CHOPS compounds which can derive from reaction processes of organophosphates with sulfur species and are likely of anthropogenic origin (Chen et al., 2023). Their origin remains unknown and needs further investigation.

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The distinct chemical compositions observed across our samples from different altitudes highlight the complexity of atmospheric dynamics and processes. Additionally to the chemical composition, the size distribution highlights the differences between ground pass and UTLS region, showing lower particle mass and a larger contribution of particles between 500 and 1000 nm in the UTLS (Fig. S6). The atmospheric composition is influenced by both anthropogenic and biogenic sources and il-
255 lustrates different transport processes. Our results particularly highlight the role of altitude and geographical location in relation to the chemical composition of the organic aerosols.

3.4 Origin of the air masses and chemical composition

As the origin of air mass plays a decisive role for the occurring chemical composition of the atmosphere, we calculated 10-days backward trajectories. The trajectories are classified according to the atmospheric region at their start position according to our
260 analysis in section 3.3. Additionally, it has to be noted that the kinematic trajectories based on ICON-global data only capture large-scale ascent, but not vertical motion represented in ICON-global by a convection parameterization. We therefore combined the trajectory position with high-resolution MSG data (IR 10.8 μm and WV 6.2 μm channels) to investigate a potential co-location of trajectories in the upper troposphere ($z > 7$ km) and convective-type clouds. Figure 4 shows a comparison of the



Table 2. Relative abundance in % of the elemental composition groups CHOn, CHOa, CHNO, CHOS, CHNOS, CHOP, CHN and *other*, measured with Orbitrap mass spectrometry in negative ionization mode. The samples displayed correspond to the region where the filter was sampled, including the ground station at the Taunus Observatory.

Sample and altitude range	CHOn	CHOa	CHNO	CHOS	CHNOS	CHOP	CHN	other
F05 - The Fold								
Troposphere 6–7 km	35.05	1.90	4.93	7.75	3.36	17.45	0.03	29.52
Troposphere 7–10 km	55.89	3.54	4.06	21.17	10.69	0.37	0.04	4.24
Stratosphere 10 km	48.53	3.03	43.63	3.86	0.57	0.09	0.00	0.29
Troposphere 10–8 km	45.84	2.94	4.55	21.19	5.39	0.34	0.60	19.16
F09 - Convection Hunting								
Troposphere 10 km	53.91	0.69	2.20	5.78	13.69	7.12	0.03	16.58
Ground Pass 0.9 km	74.19	3.82	4.48	6.66	2.59	2.29	0.03	5.93
Troposphere 8–12 km	48.15	1.88	3.34	4.17	2.07	9.91	0.21	30.26
Stratosphere 12 km	43.42	1.62	3.86	16.82	7.07	7.32	0.01	19.88
Ground Station 0.8 km	83.77	4.02	1.76	5.30	4.58	0.28	0.00	0.28

path between stratospheric and tropospheric air masses for F05 and F09. The displayed trajectories as well as the corresponding chemical composition displayed in a pie chart, are based on F05 filter 4/ F09 filter 2 (tropospheric region) and F05 filter 3/ F09 filter 4 (stratospheric region). Trajectories of the path corresponding to the remaining filters of F05 and F09 as well as additional information on vertical lifting prior to the observations can be found in Fig. S7-S15.

The LAGRANTO trajectories, calculated from the F05 flight path during the sampling period of filter 3, originate from a large area spanning parts of Alaska, Canada, Greenland, Finland and the eastern coast of North America (Fig. 4 top left) and were transported eastwards across the Atlantic towards Europe. The air mass sampled has only a small contribution (below 5 %) of strongly rising parcels and remains at altitudes above 9 km in the ten days prior to the observations (Fig. S9), with its stratospheric origin supported by trace gas data (Fig. 2).

The trajectories for flight F09 indicate that the measured stratospheric air masses (filter 4) originate from the stratosphere over eastern Canada and the western contiguous United States (Fig. 4 top right). These air masses remain at altitudes above 10.5 km in the ten days prior to the observations (Fig. S12, bottom right) and Over the course of five days, the trajectories traverse the Atlantic Ocean, passing the southern tip of Greenland, before turning south towards the Portuguese coast and then arching north again towards the flight area. Their flow pattern, altitude and longitude range between 40 and 60 °N suggest that the air masses are transported by the polar jet stream. Compounds contained in this air mass may derive from local pollution events such as biomass burning events that became globally distributed by long range transport in the jet stream (Stohl, 2001; Dicker-

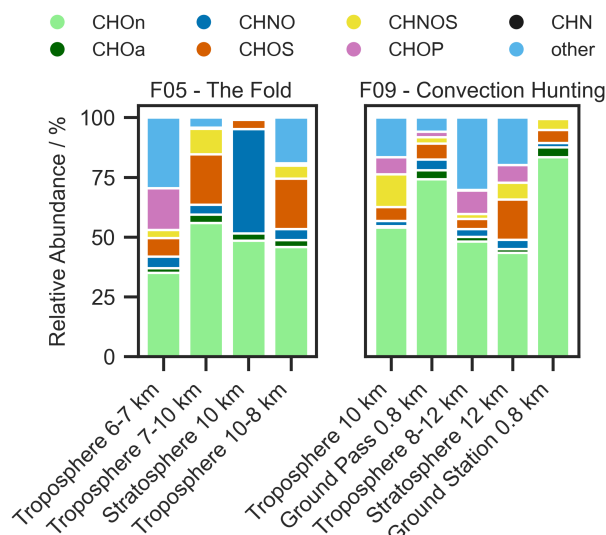


Figure 3. Relative abundance in % of the elemental composition groups CHOn, CHOA, CHNO, CHOS, CHNOS, CHOP, CHN and *other*, measured with Orbitrap mass spectrometry in negative ionization mode (also given in Table 2). Displayed are the filters that were sampled throughout the flights F05 and F09. In panel F09, the filter sampled at the ground station Taunus Observatory is added.

son et al., 2007). Notably, transport from North America to Europe, as well as from Canada to Europe, has been described in several studies (Forster et al., 2001; Joppe et al., 2024; Huntrieser et al., 2005; McKendry et al., 2001).

However, they are co-located with a convective system over the French Channel coast in the early hours of 18th June (4-7 UTC, Fig. S15 top left, green contours) and convection over Northern Germany from about 14 UTC (Fig. S15 bottom left, green contours). The air mass altitude at these times is around 12 km. Therefore, there is a possibility of convective transport, mainly from overshoots or turbulent mixing at the top of the anvil, influencing these air masses and possibly transporting organic aerosol from the ground into the stratosphere.

The trajectories calculated for the tropospheric air samples measured during F05 (filter 4) originate from a large area covering the western North Atlantic Ocean and the northern United States and have altitudes below 2 km mainly over the oceanic region (Fig. 4 bottom left and Fig. S11 bottom right). This supports the trace gas observations indicating air of tropospheric origin (Fig. 2). The air masses feature a warm conveyor belt-like ascent over the western North Atlantic Ocean (Fig. S11 bottom left) at times around four to seven days prior to the arrival in the measurement region. Strongly ascending air masses constitute about 25 % of the sampled air mass (Fig. S10). This suggests that their transport might be facilitated by low scale transport and uplift that occurred before the flight.

The trajectories calculated for the tropospheric air samples collected during F09 (filter 2) originate from a smaller area than those for F05, covering the North Atlantic Ocean and the north-west of Africa (Fig. 4 bottom right). The air masses are of



lower-tropospheric origin and mainly found at altitudes below 4 km in the ten days prior to the observations (Fig. S12 top right), covering the planetary boundary layer and the free troposphere. Trajectory positions below 2 km are spread widely over South-western Europe as well as the central North Atlantic (Fig. S14 top right). Compounds within these air masses can be transported on intercontinental scale, which is likely supported by strong winds and a mostly stable environment (Zhuang et al., 2018).

Using backward trajectories calculated by LAGRANTO, we were able to support the stratospheric or tropospheric origin of air samples collected during flights F05 and F09, which helps us investigate the correlations of atmospheric origin and anthropogenic organic compounds. Further, they highlight varying influences of organic compounds ranging from intercontinental upper tropospheric transport, deep convection and warm conveyor belt uplift. However, a larger number of flights would be required to further investigate the links between origin and anthropogenic organic compounds, and we strongly emphasize the urgency of further research into these links.

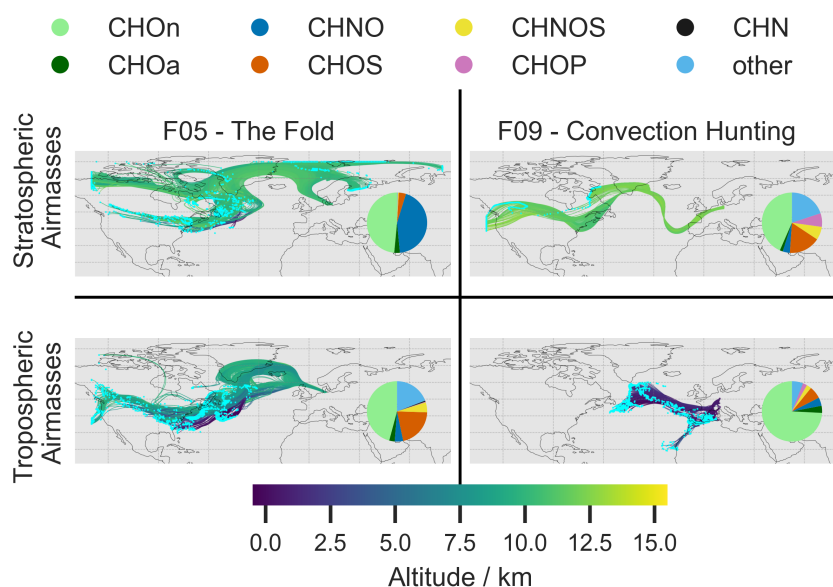


Figure 4. Five-day backward trajectories of the sampled period for F05: filter 3 (stratospheric) and 4 (tropospheric) and F09: filter 2 (tropospheric) and filter 4 (stratospheric). The altitude of the air mass is displayed in km. The origin of the respective air mass is marked by a blue dot.

3.5 Non-target analysis of organic compounds within different parts of the UTLS

To understand the distribution of organic compounds throughout the UTLS and figure out possible transport processes, we investigated all compounds depending on their occurrence in either stratospheric and/ or tropospheric air masses (Fig. 5) and

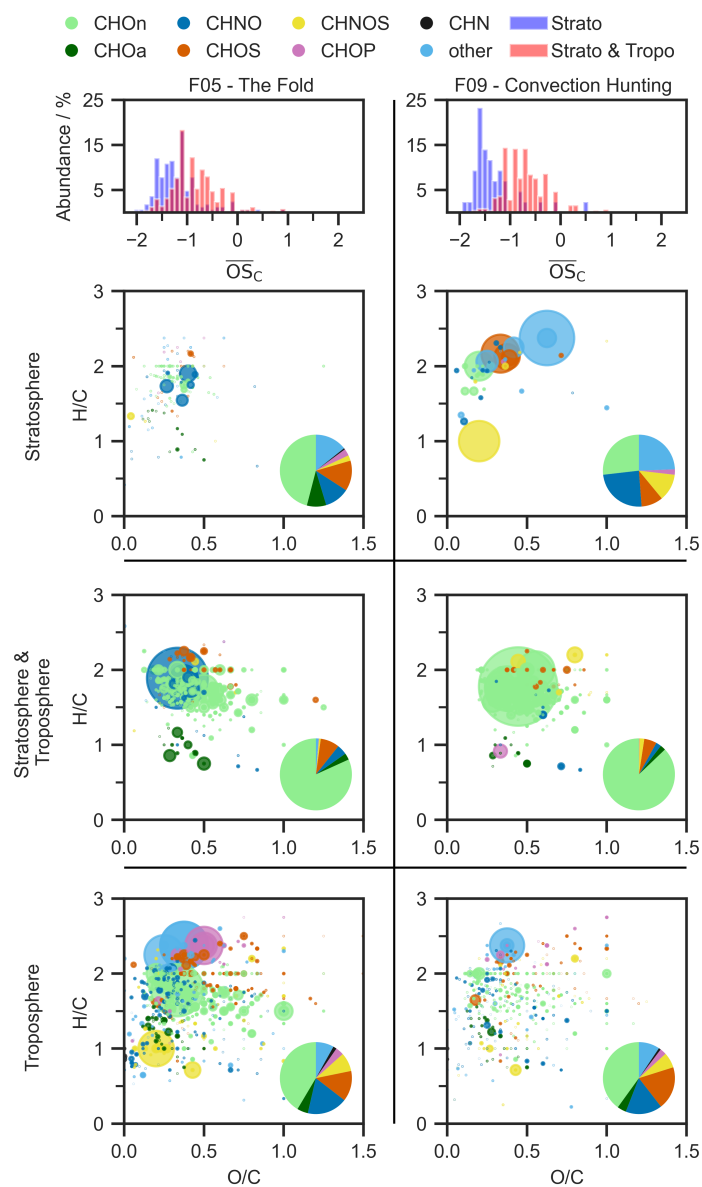


Figure 5. The uppermost panel displays the distribution of average oxidation state of the carbon atom ($\overline{OS}_C$) of all compounds for aerosols found in the stratosphere and aerosols found in stratosphere and troposphere. Below are three Van Krevelen plots, which show aerosols that are either found only in the stratosphere, the stratosphere and troposphere or only the troposphere. The y-axis displays the H/C ratio and the x-axis the O/C ratio. The circle diameter of each compound represents the relative intensity, and the circle colour the group that is assigned (CHOn, CHOA, CHNO, CHOS, CHNOS, CHOP, CHN and other). The pie chart contains the relative number of each compound group, contributing to the total compounds displayed. The left-hand side is F05 and the right-hand side is F09. All compounds displayed were measured in HESI negative ionization mode.



present novel insights on a molecular level. There are only a few studies concerning the general distribution of aerosol between troposphere and stratosphere, and hence there are no characteristics of aerosols studied so far that explain the different occurrences.

For our analysis, compounds on the stratospheric filter with a signal intensity one thousand times higher than on all other filters of the respective flight (stratospheric filter: F05 filter 3 and F09 filter 4) are assigned uniquely to the stratosphere. Compounds that can be found in the stratosphere and on every other filter of the corresponding flight representing the troposphere are assigned to both regions. The compounds that cannot be found on stratospheric filters are assigned uniquely to the troposphere. All compounds are displayed in a Van Krevelen diagram, which shows the O/C against H/C ratio, indicating degree of saturation and oxidation state. The pie charts indicate the relative number contribution of each elemental composition group. Additionally, an overall feature of the compounds, which is the average oxidation state of the carbon atom ($\overline{OS}_C$), is compared to obtain deeper insights into the compounds compositional differences.

Comparison of chemical characteristics

Starting with the difference in characteristics of stratospheric aerosols or aerosols that can be found in both regions, we investigated the $\overline{OS}_C$ of every compound against its respective carbon atom number. Overall, the $\overline{OS}_C$ of the compounds shown ranges from -2 to $+1$, which is the range where most organic aerosols appear. As described by Kroll et al. (2011), the formation of C–O bonds or cleavage of C–H bonds leads to an increase in $\overline{OS}_C$, such that more oxidized species exhibit a higher oxidation state. Notably, compounds that are unique to the stratospheric air for both flights tend to have lower $\overline{OS}_C$, indicating less oxidized species, an observation that has also been reported from balloon-borne measurements in the stratosphere (Benoit et al., 2023). Since the stratosphere contains less water vapor than the troposphere and thus a decreased production of OH, the oxidative capacity differs for each region within the UTLS and is lower in the stratosphere than in the troposphere (Esler et al., 2001). The differences are more pronounced for the convection hunting flight than for the fold, where the overlap of both compound groups is larger, but still showing the tendency of stratospheric compounds appearing in the lower $\overline{OS}_C$.

Compounds unique to the troposphere

First, we want to examine the compounds exclusively detected in the troposphere. Since they cannot be found in the stratospheric samples, they do not seem to undergo vertical transport processes neither upwards nor downwards from the stratosphere, but might experience atmospheric processing. The compounds described can rapidly deplete by reaction with OH and nitrate radicals (NO_3), thus forming new compounds (Eliason et al., 2004; Ellison et al., 1999). However, if appearing in the particle phase, semi-volatile organic compounds can be shielded against atmospheric oxidation, thus enhancing their atmospheric lifetime up to a week (Socorro et al., 2017). We found 604 compounds which are exclusively found in F05's tropospheric samples and 989 compounds that are found in F09. Here, we have an overlap of 194 compounds that are present in both flights. The contribution of different compound groups is almost identical in both flights and shows great diversity, with CHOn predominating over CHNO and CHOS. In both flights we found several oxidation products of anthropogenic and biogenic SOA that lead to a broad variety of compounds and are commonly reported within the boundary layer (Hallquist et al., 2009; McFiggans et al.,



2019). Amongst those we identified biogenic SOA like suberic acid and ozonolysis oxidation products of α - and β -pinene, limonene and β -caryophyllene that match the *Aerosolomics* database ($C_8H_{14}O_4$, level 2; $C_8H_{12}O_6$, level 2; $C_9H_{14}O_4$, level 2; $C_8H_{14}O_5$, $C_5H_6O_4$, level 2; $C_6H_{10}O_4$, level 1).

Another important source of atmospheric organic aerosol is biomass burning, which can be traced by a variety of organic compounds (Wang et al., 2017; Simoneit and Elias, 2001). Amongst those, we identified phthalic acid, nitrosalicylic acid and nitrophenol ($C_8H_6O_4$, level 1; $C_7H_5NO_5$, level 3; $C_6H_5NO_3$, level 1) (Wang et al., 2017). Likewise, organosulfates are known to contribute to a significant fraction of organic particulate matter, occurring in a large variety (Surratt et al., 2008; Tolocka and Turpin, 2012; Ma et al., 2025). The organosulfates appearing in the area $H/C > 2$ show an aliphatic character and likely originate from combustion of sulfur containing fossil fuel and ship emissions (Blair et al., 2016; Qi et al., 2021). We found a prominent related compound, which is a carbonyl hydroperoxide that can further react to its dihydroperoxide and subsequently forms an organosulfate ($C_6H_{16}O_3$, level 4; $C_9H_{16}O_5$, level 4; $C_9H_{17}O_7S$, level 4) under acidic conditions (Riva et al., 2016).

Notably, F05 shows some large signals that are not as dominant in F09 respectively, which were attributed to *other*, CHOP and CHNOS. They were identified as $C_8H_{19}O_3PS$, level 4; $C_8H_{19}O_4P$, level 4; $C_{12}H_{27}O_4P$, level 1 and $C_5H_{11}NO_4S$, level 4, with a suspected anthropogenic influence, as some organophosphates are used as flame retardants (de Boer and Stapleton, 2019).

Compounds found in the troposphere and stratosphere

Looking at compounds that are found within the troposphere and the stratosphere, a pronounced change in chemical composition of compounds contributing to this group is notable (Fig. 5). Here, over 75 % of the compounds are attributed to the CHOn group, followed again by CHNO and CHOS. 513 compounds are found for F05 and 502 for F09 with an overlap of 290 compounds that appear in both flights. For this compound category, the similarities between both flights seem to be larger than for the tropospheric-only compounds, emphasizing a distinct characteristic that allows them to cross the tropopause. Moreover, their occurrence in both regions underlines the previously discussed mixing processes in the UTLS taking place (Holton et al., 1995; Shapiro, 1980).

While the H/C ratio for compounds in the troposphere covers a broad range, most compounds found in both regions have a H/C ratio between 1.5 and 2.2, suggesting a rather aliphatic character. However, the overall range of O/C ratio is the same, with the majority appearing around 0.2–0.6. Moreover, a fraction of organosulfates appearing in the region of $H/C > 2$ remains, while the prominent organophosphates and compounds in *other* disappear.

In contrast to F09, F05 shows a strong CHNO signal which we attributed to $C_9H_{17}NO_3$, level 4. This compound was already reported by Leppla et al. (2025) and Kourtchev et al. (2016) in the context of biomass burning chamber experiments and field measurements in the Amazon rainforest. According to the trajectories source region, Canadian wild fires during that time might provide an explanation for the compounds observed (Filonchik et al., 2025). Other compounds are traced back to oxidation products of biogenic as well as anthropogenic VOC. Notably, we found several oxidation products driven by OH radicals, which leads to different products than ozonolysis, which were dominantly found in the troposphere. The OH concentration depends on different formation mechanisms that eventually lead to an increased concentration in the sampled region (Crutzen and Zimmermann, 1991). Especially due to varying residence times in the air parcel, an increased exposure to OH might ex-



plain the observed compounds (Lelieveld et al., 2016). The lifetime of atmospheric products can vary from minutes to hours depending on the precursor (e.g. sesquiterpene vs. monoterpene) and whether they are oxidized by OH or O₃ (Atkinson and Arey, 2003). We were able to identify the photooxidation products of the biogenic VOC β -pinene, β -caryophyllene and δ -3-carene (C₅H₈O₄, level 2; C₉H₁₄O₃, level 2; C₉H₈O₄, level 2). Additionally, a photooxidation product of the anthropogenic VOC 1,2,4-trimethylbenzene was identified (C₉H₈O₄, level 2). Moreover, several dicarboxylic acids from biogenic origin like levulinic and azelaic acid as well as from anthropogenic origin like adipic acid were found in both troposphere and stratosphere (C₅H₈O₃, level 1; C₉H₁₆O₄, level 1; C₆H₁₀O₄, level 1) (Mochida et al., 2003; Hansen et al., 2014). Commonly reported sources for these compounds are fossil fuel combustion (Kawamura and Kaplan, 1987), meat cooking (Rogge et al., 1991; Schauer et al., 1999) and biomass burning (Lefer et al., 1994). Dicarboxylic acids hereby have a low water vapor pressure and make up a large fraction of water-soluble organic aerosols due to their high water solubility (Saxena and Hildemann, 1996). Because of their physico-chemical characteristics, they are able to affect the properties of atmospheric particles including size, hygroscopy and activity as cloud condensation nuclei (Lightstone et al., 2000; Jacobson et al., 2000; Xu and Zhang, 2012). Similarly in F09, we identified azelaic, adipic acid, and photooxidation products of several biogenic VOC, such as β and α -pinene, two products of β -caryophyllene and δ -3-carene as well as the anthropogenic VOC naphthalene (C₈H₁₂O₄, level 2; C₉H₁₄O₄, level 2; C₉H₁₄O₅, level 2; C₉H₁₄O₄, level 2; C₉H₁₄O₄, level 2; C₆H₁₀O₄, level 2; C₈H₁₂O₄, level 2).

Compounds unique to the stratosphere

Compared to the molecular fingerprint observed in the troposphere or for both regions, the compounds unique to the stratosphere show a completely different chemical profile. These compounds might not be transported vertically, but horizontally due to long-range transport or formed by atmospheric processes such as oxidation reactions or condensation of volatile species (Stohl, 2001; Dickerson et al., 2007).

Here, only a few compounds are found to be unique to the stratosphere, which are 166 compounds for F05 and with 43 even less for F09. The difference in number of emerging unique stratospheric compounds can be explained by differing meteorological scenarios, where F09 captures convective processes that increases the probability for vertical transport and mixing. As a consequence of that it is possible that more compounds are found in both layers rather than being identified as unique to the stratosphere. Surprisingly, there is no overlap between the compounds of the two flights, emphasizing the chemical variability of stratospheric organic aerosol composition. Even though a comparably small number of compounds is found as stratospherically unique, especially those could be of large interest to understand their origin, atmospheric mixing, transport processes and the anthropogenic influence on stratospheric OA.

Outstanding signals in F09 are organothiophosphates and a CHNOS compound (C₈H₁₉O₅PS₂, level 3; C₁₂H₂₇O₅PS₂, level 3; C₁₅H₁₅NO₃S₂, level 4). These compounds were not found to be reported in other studies and need to be studied further, especially regarding their effects within the stratosphere. Notably, C₁₅H₁₅NO₃S₂ has an H/C ratio close to 1, indicating an aromatic compound. In this regard, studies by Li et al. (2020) point out the light-absorbing potential which should be noted especially in context of the stratosphere.

While in F05 the compound group CHOn is dominant, there is an almost equal distribution between CHOn, CHNO and *other*



for F09. Both flights show additionally organosulfates, which show again an aliphatic character (e.g. $C_{18}H_{38}O_7S$, level 4; $C_{16}H_{34}O_6S$, level 4; $C_{12}H_{26}O_5S$, level 4; $C_{14}H_{28}O_5S$, level 4). Some pronounced CHNO signals in F05 show a homologue series of organonitrates that likely derive from oxidation chemistry by NO_3 radicals and biogenic derived monoterpenes (Curtius et al., 2024) ($C_9H_{15}NO_3$, level 4; $C_{10}H_{19}NO_4$, level 4; $C_{11}H_{17}NO_4$, level 4; $C_{12}H_{21}NO_5$, level 4). Additionally, CHNO compounds with a high nitrogen number, such as $C_{15}H_{26}N_6O_4$, level 4 were found, leading to an O/N ratio below 3 (Fig. S6). These compounds indicate the oxidation of nitrogen containing precursors such as amides, that are derived from anthropogenic sources like vehicular exhaust (Yang et al., 2022), rather than organic compounds that react with NO_3 , attaching a nitrate group (Mayorga et al., 2021).

To conclude this comparison, we want to highlight the novel approach of analyzing different regions of the atmosphere via non-target analysis. By this, we were able to highlight differences in chemical patterns and characteristics, that occur for each region. We saw a broad variety of compounds for both troposphere and stratosphere, differing most for the stratosphere between both flights. Also, indications for transport processes were highlighted, stressing the potential of this measurement method.

3.6 Identified anthropogenic influence in the UTLS

To highlight the possibilities of a targeted approach and focus more on investigating the anthropogenic influence within the UTLS, we also searched for target analytes with known origin from anthropogenic emissions. First, we focused on an approximation of the general anthropogenic influence, calculated by the ratio of azelaic acid, a biogenic marker and adipic acid, an anthropogenic marker, as suggested by Ho et al. (2006). Both substances were identified on the confidence level 1 by using authentic standards.

Notably, the strength of biogenic sources varies significantly (Fig. S17) for the different flights as well as the different altitudes. This emphasizes the present anthropogenic influence and complexity of formation and transport processes, which is coherent with the chemical composition we observed. While both stratospheric samples show almost the same ratio of biogenic to anthropogenic source, the remaining samples go from predominantly anthropogenic influence in the upper troposphere to vast biogenic influence and back to largely anthropogenic influence. The non-existing trends stress the urge for more samples to gain a comprehensive understanding of the different influences.

As for target analytes, we want to give a first impression of compounds whose role and existence in the UTLS is largely unknown and yet are found via our UHPLC-HRMS setup and offline filter sampling. Moreover, target analytes and especially anthropogenic pollutants contain valuable information about redistribution processes from ground to the UTLS that needs to be studied further.

Here, we focused on organophosphates which are or were used as flame retardants, lubrication oil, that is applied in civil aviation, and PFAS, which are known as the "forever chemicals", with a lifetime of decades, centuries or eventually longer (Brunn et al., 2023). The long lifetime of these pollutants in the free troposphere extends their impact from the regional to the continental or even global scale distributed by long-range transport. For this purpose, we investigated the compounds based on the exact masses as well as the isotopic fine structure (Fig. S18) and aim for deeper investigations and comparison with authentic standards.

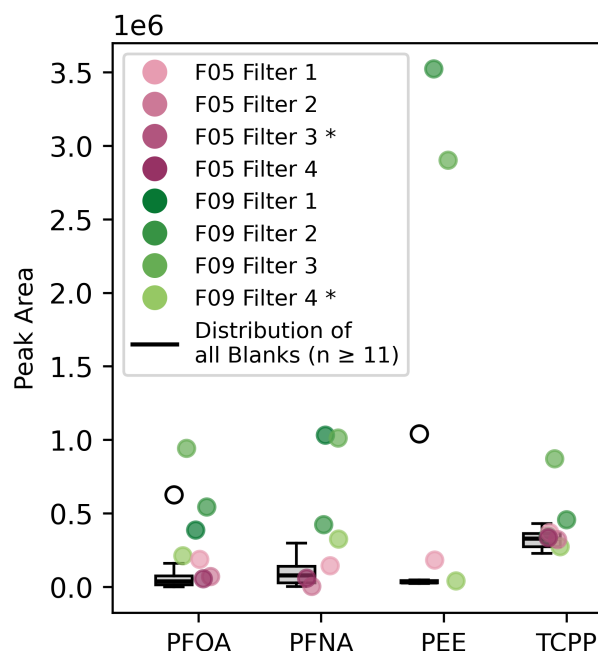


Figure 6. The colored dots display the peak area (y-axis) of the target compounds PFOA, PFNA, PEE and TCPP for the respective filter. The boxplot next to it displays all filter blanks ($n \geq 11$) that have been collected throughout the whole campaign. The stratospheric filters are marked with a star (*). Colorless circles mark outliers that are defined as $x < Q1 - 1.5 \cdot IQR$ or $x > Q3 + 1.5 \cdot IQR$ with $IQR = Q3 - Q1$, with IQR as the interquartile range.

The selected pollutants perfluorooctanoic acid (PFOA), perfluorononanoic acid (PFNA), Tris(2-chloropropyl) phosphate (TCPP), and the pentaerythritol ester (PEE, $C_{29}H_{52}O_8$) were especially found in F09, which represents a flight over land and suggests redistribution from emission sources on ground (Fig. 6). In terms of the pentaerythritol ester ($C_{29}H_{52}O_8$), which is used as lubrication oil in jet engines (Ungeheuer et al., 2021), the tendency to form new particles is already studied (Ungeheuer et al., 2022). Here, we only found the major PEE and were not able to measure all other esters that occur in the lubrication oil. The ground pass, which exhibits the largest signals for PEE, took place relatively close to the Frankfurt airport, though we did not specifically aim for measuring aircraft emissions. That being the case, their occurrence within the UTLS is an important and understudied source of anthropogenic organic aerosols (Decker et al., 2024).



460 4 Conclusions

In this study, we demonstrated the successful application of UHPLC coupled with HRMS measurements and a subsequent non-target analysis to investigate organic aerosols within the UTLS. By collecting filters with our in-house built multi-channel offline filter sampler SOAP during eight research flights, we were able to characterize the chemical composition of different parts of the UTLS. The eight flights captured various meteorological events with flight tracks over land and sea, reaching
465 altitudes of up to 12 km. Our focus on a tropopause fold and convective systems allowed us to examine potential transport mechanisms for different aerosols, specifically highlighting F05 and F09 as case studies for this and further studies focusing on more detail on likely transport mechanisms.

Equipped with a comprehensive set of instruments, we were able to distinguish between tropospheric and stratospheric air masses by examining trace gases, capturing and separating their aerosol compositions on different filters. A non-target anal-
470 ysis revealed significant variations between samples from different regions within the UTLS. In particular, we observed a notable abundance of CHNO compounds in the stratosphere over the sea, while above land, a broader chemical variety prevailed. The flights closer to the ground aligned with our expectations of chemical composition and were comparable to ground measurements taken at the Taunus Observatory at Kleiner Feldberg, contrasting with the more diverse chemical profile of the stratosphere.

475 There is currently a lack of studies concerning the general distribution of aerosols between the troposphere and stratosphere, which limits our understanding of the characteristics that explain their differing occurrences. The origin of the air masses determined by LAGRANTO calculations suggests long-range transport of stratospheric air masses, possibly influenced by the polar jet stream, as well as medium-range transport in the troposphere and planetary boundary layer. In addition, the kinematic trajectories indicate strongly varying contributions from large-scale ascent and convectively transport including overshoots and
480 turbulent mixing in the UTLS, emphasizing the complexity of aerosol found in the UTLS. Most air masses originated over North America and the Northern Atlantic Ocean.

A deeper investigation of compounds unique to either the troposphere or stratosphere revealed changes in general chemical properties. We noted a decrease in the average oxidation state of carbon in compounds found only in the stratosphere. Furthermore, we identified dicarboxylic acids, anthropogenic, and biogenic VOC oxidation products, as well as organosulfates in both
485 regions, suggesting redistribution from ground emissions. An additional target analysis of the anthropogenic pollutants TCPP, PFOA, PFNA, and a PEE used as jet oil highlighted the extensive capabilities of our measurement technique and underscored the anthropogenic influence detected in the stratosphere, which warrants further investigation.

Our analysis is presently limited by the sampling duration, which affects detection limits, and our analytical setup restricts the range of detectable compounds via HESI. However, we identified specific compounds such as $C_8H_{19}O_5PS_2$ and $C_{15}H_{15}NO_3S_2$
490 that, with future research, could yield valuable insights into various atmospheric processes. Our method, combining offline filter sampling with UHPLC-HRMS measurements and non-target analysis, grants novel insights into the chemical composition of the UTLS and opens new avenues of inquiry to advance our understanding of mixing processes.

This study confirms the anthropogenic influence in the UTLS region, providing a closer examination of potential reaction and



transport pathways. Further research is essential to assess the implications how small-scale and synoptic-scale transport redis-
tribute anthropogenic aerosol throughout the atmosphere. Understanding changes in the number and mass of particles in the
UTLS resulting from anthropogenic activities is crucial. Overall, our results contribute to a broader understanding of the chem-
ical composition in the UTLS region and emphasize the necessity for additional studies utilizing advanced analytical methods
like HRMS to unravel the complexities of atmospheric interactions. We provide evidence of a significant anthropogenic foot-
print in the UTLS aerosol, offering essential insights to enhance our comprehension of climate-relevant processes and improve
future atmospheric modeling efforts aimed at predicting the behavior of aerosols and their implications for air quality and
climate change.

Data availability. The data is available from the Zenodo online repository <https://doi.org/10.5281/zenodo.15680610>. MSG satellite data is available from the EUMETSat archive.

Author contributions. AB collected and measured the samples, analyzed the data and wrote the manuscript. AV and TH planned and sup-
ported during the campaign. FU helped performing the target analysis. PJ, JS, JW are responsible for the AMS and UHSAS data. NE, IK,
HB and PH provided CO, N₂O and O₃ data. RR, LM, TK, KD, AK, PW and SH were responsible for planning, manufacturing, certifying
and installing SOAP. AM and CS calculated and provided the backward trajectories for the respective flights. PJ and SR acted as scientific
operators during the flights. We acknowledge the DWD open data server, which was used to retrieve operational data during the campaign.
EuMeSaT Meteosat Second Generation HRSEVIRI data shown in Fig. S15 has been downloaded via EUMDAC in May 2025. All authors
contributed to and commented on the manuscript.

Competing interests. The authors declare that they have no conflict of interest.

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