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Investigating the evolution of water-soluble organic carbon in evaporating cloud water†

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Cloud cycling plays a key role in the evolution of atmospheric particles and gases, producing secondary aerosol mass and transforming the optical properties and impacts of aerosols globally. In this study, bulk cloud water samples collected at Whiteface Mountain (Wilmington, NY) in the summer of 2017 were aerosolized, dried to 50% RH, and analyzed for the evaporative loss of water soluble organic carbon (WSOC) and for brown carbon (BrC) formation. Systematic WSOC evaporation occurred in all cloud water samples, while no evidence for drying induced BrC formation was observed. On average, 11% ($\pm 3\%$) of WSOC evaporated when the aerosolized cloud droplets were dried to 50% RH, though this represents a lower bound on the WSOC reversibly partitioned to clouds due to experimental constraints. To our knowledge, this represents the first direct measurements of organic evaporation from actual cloud water undergoing drying. Formate and acetate contributed 19%, on average, to the evaporated WSOC, while no oxalate evaporation occurred. GECKO-A model simulations were carried out to predict the production of WSOC compounds that reversibly partition to cloud water from photooxidation of an array of VOCs. The model results suggest that precursor VOC identity and oxidation regime (VOC:NO_x) have a dramatic effect on the reversible partitioning of WSOC to cloud water and the abundance of aqSOA precursors, though the higher abundance of reversibly partitioned WSOC predicted by the model may be due to aqueous production of low-volatility material in the actual cloud samples. This study underscores the importance of the large fraction of unidentified compounds that contribute to WSOC in cloud water and their aqueous processing.

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Environmental significance

Clouds serve as important reservoirs for the transformation of organics in Earth's atmosphere. Water-soluble gases can partition to cloud droplets and react to form lower-volatility compounds that increase the SOA burden. Such aqueous reactions can also form light-absorbing brown carbon (BrC), a process accelerated when droplets evaporate. Despite their potential importance for global aerosol concentrations and radiative effects, there are immense challenges in directly measuring in-cloud transformations of organics *in situ*. Herein, we aerosolized and dried cloud water samples collected at Whiteface Mountain (NY) to mimic a cloud cycle. All samples showed WSOC evaporation with drying but negligible BrC production. The results advance our understanding of in-cloud organic chemistry and provide a critical link between prior laboratory and field observations.

1. Introduction

Low-level clouds are abundant in Earth's atmosphere and interact with boundary layer gases and particles to affect the fate and transport of trace species. Cloud processing alters the size

distribution and chemical composition of ambient aerosol.¹ Photooxidation of anthropogenic and biogenic volatile organic compounds (VOCs) forms water-soluble organic compounds (WSOC) commonly observed in clouds and fogs.² Partitioned species undergo chemical transformation in the aqueous phase that is often different from gas-phase oxidation pathways, and can favor the formation of low volatility compounds that contribute to secondary organic aerosol (aqSOA) mass.¹ Cloud radiative impacts are also substantial and include potential production of brown carbon (BrC), a short-lived climate forcer, aloft.³ While ubiquitous and important, large uncertainties in cloud physical predictions and aqueous-phase chemical mechanisms hinder accurate quantitative understanding of

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aqSOA contributions to tropospheric aerosol burdens, though it is expected to be substantial.^{4,5}

Ambient cloud physics and cloud microphysics are routinely studied in field and laboratory experiments,⁶ but the investigation of cloud chemistry is not as widespread. The majority of liquid water droplets (>90%) evaporate after air parcels pass through clouds.⁷ The fundamental chemistry and potential WSOC evaporative loss during drying of ambient cloud droplets remain poorly understood, as does BrC formation during cloud droplet drying. Laboratory studies of organic partitioning during droplet evaporation investigate individual compounds^{8–11} and find that observed gas- and condensed-phase concentrations can deviate from thermodynamic partitioning predictions up to several orders of magnitude, especially for more polar species.¹² This inhibits the development of chemical mechanisms to describe cloud processing of complex organic mixtures given that single component systems are difficult to predict. Evaporative formation of BrC is observed in laboratory experiments with simplified droplet and particle compositions, but not in the drying of ambient aerosols sampled in the eastern US.^{11,13,14} Such conflicting results challenge efforts to develop parameterizations that can be used in models to evaluate this process on larger scales. During a comprehensive and nearly ideal Lagrangian study of cloud chemistry at Schmücke Mountain, it was difficult to constrain physical losses (droplet impaction on vegetation), and chemical transformations from concurrent upwind, in-cloud, and downwind measurements, precluding quantitative assessment of gas-aqueous partitioning for most organic compounds.¹⁵ Further, the majority of WSOC in clouds and fogs are unidentified on a molecular basis,² so the source of these compounds (whether present in the activated CCN or partitioned from the gas phase upon cloud droplet formation) and their subsequent fate under cloud droplet evaporation remain uncharacterized.¹⁶

In this study, we evaluate the cloud processing of WSOC by analyzing cloud water samples collected at Whiteface Mountain, Wilmington, NY, a remote forested site influenced by air-masses from the Central Midwest, Great Lakes and Canada.^{17–19} We characterize the evaporative loss of WSOC and formation of BrC in aerosolized cloud water under conditions of drying. We also model the equilibrium partitioning of WSOC formed from the photooxidation of various precursor VOCs under high- and low-NO_x conditions with the GECKO-A box model.

2. Methods

2.1 Cloud water sampling

Whiteface Mountain (WFM) is a peak in the Adirondack Mountain Range in northern New York. The summit (latitude N 44°21'58" and longitude W 73°54'10") is situated above tree line at 1483 m a.s.l., and is isolated from other peaks.¹⁷ Clouds are regularly intercepted (25–60%) at the summit in summertime.¹⁶ WFM is mostly affected by air-masses transported from the western and central United States, Great Lakes region and Canada, suggesting the organics (gas and aerosol) are aged and often dominated by contributions derived from precursor biogenic species.^{17–20} Aged biomass burning smoke intercepts

the summit of WFM and episodically influences cloud water composition.^{20,21} The Atmospheric Sciences Research Center (ASRC) at the University at Albany, Albany, NY, operates a research observatory at the summit, where cloud water sampling has been conducted regularly since 1994, historically focusing on acid deposition.¹⁷ The cloud water sampling is usually conducted in summertime (June–September) due to prohibitive wintertime weather, while gas phase measurements are performed year round.

Cloud water samples were collected during August 2017 as a part of the Cloud Processing of Organics within Clouds (CPOC) pilot study.²⁰ Cloud water was collected using a Möhnen omni-directional passive cloud water collector,²² which was deployed when the following conditions were met: (1) liquid water content measured by the Gerber Particle Volume Meter (Gerber PVM-100)²³ exceeded 0.05 g m⁻³, (2) wind speed exceeded 2 m s⁻¹, (3) temperature exceeded 2 °C (to avoid freezing), and (4) no rain was detected by a CAPMoN (Canadian Air and Precipitation Monitoring Network) resistive heating element rain sensor. The collected samples were frozen and stored in a deep freezer (–20 °C), and thawed immediately before analysis. The cloud water samples analyzed in this study were collected on August 12 (06:00–09:00), August 13 (06:00–09:00), August 13 (10:00–11:00), August 18 (06:00–09:00) and August 18 (11:00–12:00), where the time specified is local time (UTC – 4 h). *In situ* measurements of below-cloud aerosol composition on August 18 were concluded to have been urban-influenced, with an organic mass fraction of 75%, which was slightly lower than average for the CPOC pilot study.²⁰ On these 3 days (August 12, 13 and 18), cloud water samples are estimated to have been collected at 300, 490 and 400 m above cloud base, respectively, corresponding to 2.5–4 minutes of time in cloud assuming 2 m s⁻¹ updraft velocity.

2.2 Cloud water experiments and analysis

Cloud water samples were analyzed for WSOC, BrC *via* absorbance at 365 nm (Abs₃₆₅), and other inorganic and organic ions. The aim of the analyses was to (1) measure the fraction of WSOC that partitions to the gas phase upon drying, mimicking the cloud cycling effect and cloud processing, and (2) evaluate drying-induced BrC formation through the Abs₃₆₅ measurement. The experimental setup was adapted from a previous study from our group (Fig. 1).¹⁴ Cloud water was atomized (TSI, model 9302) and alternated between a dry channel, which contained a silica-gel diffusion-dryer, and a bypass channel every 15 minutes using an automated 3-way valve (Brechtel manufacturing). Note here, the cloud water samples were not exposed to any oxidants in the lab (in either the bulk or as aerosolized droplets), and no UV light source was used to initiate aqueous reactions. The drying induced BrC formation chemistry takes place due to reactants already present in the cloud water. The particles were dried to ~50% RH (not fully dried) in the dry channel, while particles in the bypass-channel experienced an RH of ~85% due to the low humidity in the makeup air. Each cloud water sample measurement was repeated 2–4 times through each channel. Particle losses in the



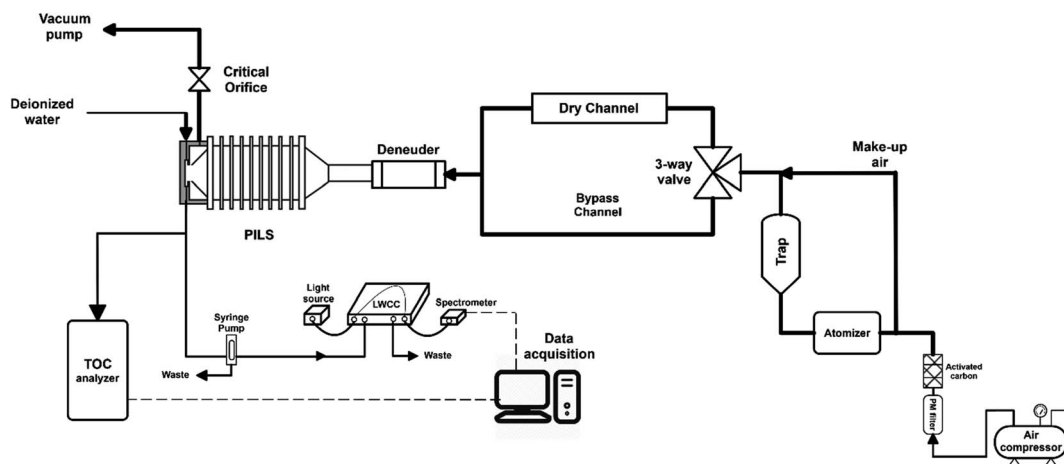


Fig. 1 Schematic of the experimental setup for laboratory analysis of the cloud water.

experimental setup were investigated with a mixed sucrose/NaCl solution. We observed no difference in particle concentrations between the two channels during the control experiments, consistent with the QA/QC results of our prior studies.^{14,24} The samples from either channel passed through an activated carbon denuder to remove VOCs prior to collection in a particle-into-liquid sampler (PILS). The PILS samples were analyzed on-line for WSOC using a total organic carbon analyzer (Sievers 900 portable, GE analytical instruments) in turbo mode (4 s resolution) and UV-absorbance using a liquid waveguide capillary cell (LWCC, model 3050, World Precision Instrument) coupled to a monochromatic 365 nm light source (Oceanoptics, LLS-LED) and a spectrometer (Oceanoptics, FLAME-S, 200–800 nm). The effective path length of the LWCC was 50 cm. The difference in WSOC and Abs_{365} from the two channels provided the measurement of drying-induced evaporation of WSOC and BrC formation, respectively. Additional samples from the PILS were analyzed off-line for inorganic (SO_4^{2-}) and organic (formate, acetate, oxalate) ions *via* ion chromatography (Metrohm 850 dual IC, Metrohm). The cation and anion columns used were Metrosep C4 – 150/4.0 and Metrosep A Supp 5 – 150/4.0, respectively. Since SO_4^{2-} is non-volatile, the SO_4^{2-} measurement through each channel was used to normalize the measurements to account for any minor differences in atomizer output during an experiment, assuming that changes in the atomizer output do not affect the relative mass concentrations of solute within the droplets produced. Since formate and acetate were not chromatographically resolved, we describe their collective behavior and concentration (together F + A) using their individual IC calibrations and the average F/A molar ratio measured in a previous study at WFM (1.9) to interpret the present results.²⁵ The uncertainty in formate, acetate and oxalate measurements were 0.15 ppbC, 0.23 ppbC and 0.1 ppbC, respectively.

Note that the RH through the bypass channel was not 100%, thus some WSOC may have evaporated from the aerosolized droplets relative to the bulk cloud water. However, the 35% RH reduction (from 85% to 50%) appears to be a reasonable drying

based on RH measurements at the WFM summit and downwind lodge site. The median RH at the summit was 98% during August 2017, consistent with the frequent presence of clouds, while the median RH at the downwind lodge site was 80% for the same period (Fig. S1†). It should be noted that the airmasses may experience both an increase in RH or further drying downwind of the lodge, so a range of dry channel RH levels should be investigated. However, limited sample volumes constrained the extent of drying RH values in this study. The drying time of ~ 7 s is sufficient for water equilibration with particles, however, organics may take longer to fully evaporate.^{8,24,26,27} In addition, further evaporation of WSOC is expected at longer drying times and lower RHs. Together, this indicates that our measurements of WSOC evaporation from the cloud water represent a lower bound on the amount of organics that may evaporate in the atmosphere during an actual cloud cycle.

2.3 Model

The photooxidation of ten atmospherically relevant organic compounds under high- and low- NO_x conditions was simulated by the Generator of Explicit Chemistry and Kinetics of Organics in the Atmosphere (GECKO-A) box model. GECKO-A models the reaction of atmospheric organic gases using laboratory-generated kinetic data or through estimates using structure-activity relationships.²⁸ The model also estimates the water solubility (Henry's constant, K_H) and vapor pressure (volatility, C^*) of thousands of oxidation products through structure-activity relationships.²⁹ For compounds whose physical constants have not been determined experimentally, the vapor pressure was estimated using the methodology in Nannoolal *et al.* (2008)³⁰ while the Henry's constant was estimated using the group contribution method for Henry's law estimate (GROMHE).¹²

The specific simulations used for this study have been described in detail before.^{29,31} Briefly, the gas-phase photooxidation of the following ten VOCs was simulated under mid-latitude tropospheric conditions: $C_{8,12,18,22}$ -alkanes, benzene, toluene, *m*-, *o*-, *p*-xylenes (xylenes considered together), C_{12} -



alkenes (with internal and terminal double bonds, considered separately), and α -pinene. The relative abundance, K_H , and C^* values of each oxidation product were determined after photo-oxidation equivalent to ~ 2 days of atmospheric aging, an appropriate value for air masses impacting WFM.^{19,32} The simulations predict hundreds or thousands of oxidation products with a wide distribution of K_H and C^* values for each VOCs considered.²⁹ Here, we extend our prior work by calculating the partitioning of each oxidation product after 2 days of gas-phase photooxidation, with no treatment of condensed-phase reactions in the aerosols or cloud water. The fraction (molar basis) of oxidation products dissolved in cloud water, ϵ_{CW} , was calculated according to Seinfeld and Pandis (2006) assuming cloud liquid water content of 1 g m^{-3} , very close to the median value for the WFM samples analyzed (1.12 g m^{-3} , Table S1†).³³ In this work, we assume liquid clouds only, to avoid complexities in partitioning that arise with ice or mixed-phase clouds.³⁴ This is consistent with summertime conditions at WFM (Table S1†). We assume equilibrium partitioning, an approach taken by most regional and global climate models, neglecting deviations that can occur due to mass transfer limitations.^{35,36}

3. Results and discussion

3.1 Experimental results and discussion

3.1.1 WSOC evaporation in cloud water droplets. WSOC evaporation was observed due to drying in all five WFM cloud water samples. On average, we observed an 11% reduction in WSOC due to evaporation, with a range of 6–13%, when dried to $\sim 50\%$ RH (Fig. 2). These results are outside experimental

uncertainty of $\pm 3\%$. In control experiments with mixed sucrose/NaCl droplets, no organic evaporation was observed, as expected for a highly soluble and nonvolatile compound. In control experiments with mixed glyoxal/NaCl droplets, approximately 17% ($\pm 3\%$) of WSOC evaporated, qualitatively consistent with previous laboratory studies involving glyoxal evaporation.^{10,14} Details of the cloud water samples and analytical results are listed in Table S1.†

To our knowledge, this represents the first direct measurements of organic evaporation from cloud water undergoing drying. In contrast, previous studies conducted on ambient aerosols in the eastern United States during summer have shown 10–30% evaporation of particulate WSOC when dried.^{14,37} The dry channel RH in the prior studies was lower (41% and 35%, respectively) than the one in the current study ($\sim 50\%$), but was subject to the same residence time of 7 s in the drying system.^{14,37} In the atmosphere, cloud droplets may undergo a greater extent of drying than aerosol depending upon the cloud properties and evolution of meteorology that may lead to additional evaporation of WSOC.³³ Future work should investigate variable drying configurations and temperatures to characterize a wider range of atmospheric conditions.

The combined contribution of formate, acetate, and oxalate to CW WSOC decreased with increasing WSOC (Fig. 3a), from a high of $\sim 35\%$ at the lowest CW WSOC (CW3, $\sim 50 \mu\text{g-C L}^{-1}$) to a low of only 7% at the highest CW WSOC (CW5, $\sim 167 \mu\text{g-C L}^{-1}$). Formic, acetic, and oxalic acids are generally the most abundant carboxylic acids found in cloud water, and the relatively high vapor pressure of formic and acetic acid indicate their likely contribution to the evaporated WSOC.^{25,38–42} Our observations suggest an array of individual compounds contributed to the evaporated WSOC, as formate and acetate (F + A) together accounted for an average of only 19% of the evaporated WSOC (C mass basis). The evaporation of F + A was highly variable across the five CW samples; we found that 0–30% of F + A was lost due to evaporation (Fig. 3b). Notably, we observed no F + A loss due to drying in CW5, the sample with the strongest

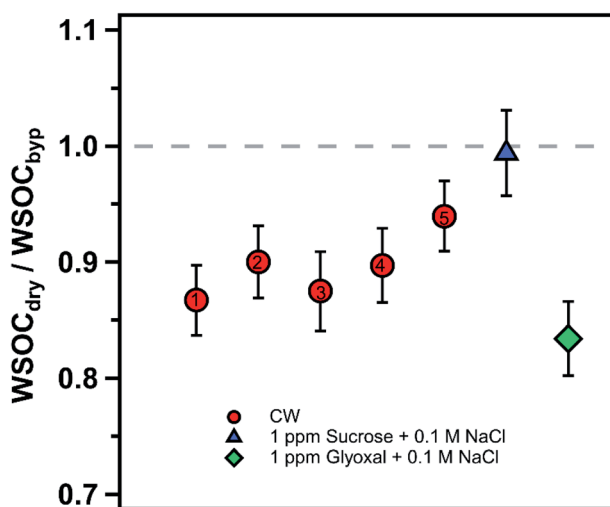


Fig. 2 Ratio of WSOC measured in the dry and bypass channels for the cloud water (CW) samples (red circles). The numbers inside the red circles correspond to the cloud water sample index (i.e., 1 denotes CW1). The error bars represent the measurement uncertainty, propagated from the accuracy of WSOC measurements. The cloud water samples are contrasted with experiments performed with non-volatile (sucrose, blue triangle) and volatile (glyoxal, green diamond) species. The dashed line at unity represents no organic evaporation due to drying, shown for visual reference.

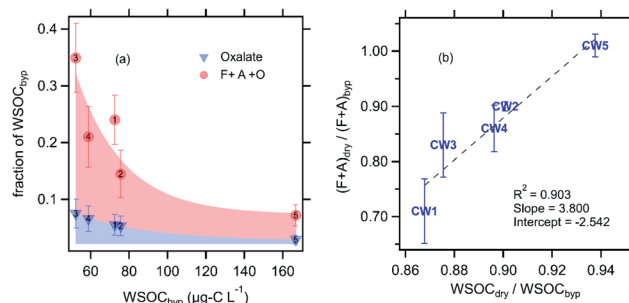


Fig. 3 (a) Contribution of oxalate (blue triangles) and the sum of oxalate, formate and acetate (red circles) to the total WSOC as a function of the WSOC concentration, all measured in the bypass channel (not dried, see Fig. 1). (b) Fraction of WSOC and F + A that evaporated with drying. Note that no oxalate evaporation was observed with drying. Error bars in each panel represent the estimated uncertainty derived from error propagation of the individual measurements. The numbers in panel 'a' correspond to the CW sample number.



signature of photochemical and aqueous-phase processing (ESI[†]). Multiple studies have shown that the gas-particle partitioning of formic and acetic acid do not typically agree with thermodynamic predictions, as the condensed-phase fraction is often higher by orders of magnitude.^{43,44} This may explain the relatively high fraction of F + A remaining after drying. We observed no oxalate evaporation in any of the cloud water samples analyzed. Fig. 3b also shows that the evaporative loss of F + A was linearly proportional to that of WSOC. While the majority of evaporated WSOC was not identified, our correlation is consistent with prior observations that F + A share similar sources and/or condensed-phase processes with this broader pool of WSOC compounds.^{45–49} For example, Millet *et al.*, (2015) showed a large concentration of formic acid in the southeastern United States, which is dominated by biogenic emissions in summertime.⁴⁶ Similarly, Paulot *et al.*, (2011) reported biogenic emissions as a major source of global formic and acetic acid concentrations.⁴⁷ As discussed previously, airmasses at WFM are expected to be predominantly influenced by biogenic emissions.²⁰ In addition, a number of studies have also established that WSOC is efficiently produced from biogenic sources, at least in summertime in the eastern United States, when the cloud water samples were collected.^{50–52}

These results demonstrate that a small but significant fraction of WSOC in clouds derives from organic gases reversibly partitioned to water. Our findings complement measurements of gas scavenging by clouds,³⁴ and direct Lagrangian sampling of gas uptake in clouds.¹⁵ These results have importance for the budget of reactive carbon in the atmosphere, and for the associated effects on air quality and climate. The reversible partitioning of WSOC to cloud water impacts the lifetime and transport of reactive organics in the atmosphere since the removal of oxygenated (and water-soluble) gases *via* dry deposition and photolysis can be substantially reduced through transfer to the aqueous phase.⁵³ The uptake of small, volatile WSOC compounds can also alter the evolution of cloud droplets and their physical properties, with implications for the climate system.⁵⁴ Volatile WSOC compounds have long been known to have the potential to form SOA through aqueous chemistry in clouds,^{2,4} and many of these compounds are semi-volatile and reversibly partition to water (*e.g.*, glyoxal, methylglyoxal).^{10,11,55} Our results are qualitatively consistent with observations from the WACS-2010 study, which suggested the formation of SOA in cloud water from dissolved organics that were too volatile to exist in the aerosol phase.⁵⁶ Although the contribution of aqueous reactions in clouds to ambient OA concentrations remains very difficult to measure or constrain, our results add to the body of work suggesting the importance of this phenomenon.¹

3.1.2 BrC formation. Fig. 4a shows Abs_{365} and WSOC measurements through the bypass channel. The cloud water Abs_{365} values are slightly higher than measurements of ambient particles (mean $Abs_{365} = 10.1 \times 10^{-3}$) in our previous study.¹⁴ Abs_{365} was moderately correlated with WSOC in the cloud water ($R^2 = 0.39$, Fig. 4a), possibly pointing to a common source. Based on the potassium (K^+) measurements in 12 hour composite samples of CW (Table S2[†]), at least three CW

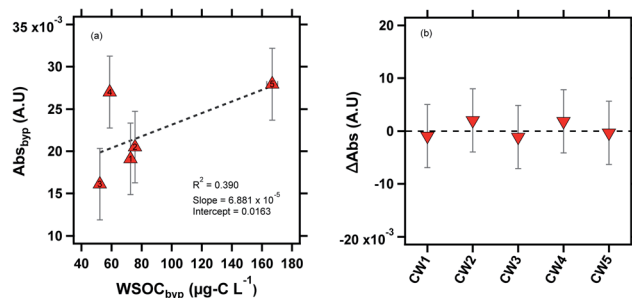


Fig. 4 Absorbance measurements of the cloud water samples at 365 nm. CW1, CW2 and CW5 were likely influenced by biomass burning, as indicated by the potassium ion concentrations (Table S2[†]). (a) Linear correlation between Abs_{byp} and $WSOC_{byp}$. (b) ΔAbs ($\Delta Abs = Abs_{dry} - Abs_{byp}$) is plotted for the cloud water samples. The subscript "byp" represents measurements through the bypass channel, and the dashed line at $\Delta Abs = 0$ line is shown for visual reference.

samples (CW1, CW2 and CW5) appear to be affected by biomass burning, the largest source of atmospheric BrC, suggesting that WSOC in these samples also derived from biomass burning. We note the limitations associated with the small sample number for this correlation analysis ($n = 5$); however, results from prior studies conducted at WFM suggest similar association between WSOC and BrC.²¹ A detailed study on cloud water samples from this campaign was performed by Lance *et al.*, (2020),²⁰ who found significant K^+ concentrations in approximately 50% of cloud water samples, and strong associations between K^+ and WSOC, indicating the regular influence of biomass burning.

Previous laboratory studies have confirmed BrC formation by aqueous reactions between dicarbonyls (glyoxal, methylglyoxal) and reduced nitrogen species (ammonium sulfate, ammonia, glycine).^{57,58} In drying particles, these reactions accelerate by several orders of magnitude due to the concentration of reactants, suggesting that secondary BrC formation in evaporating cloud droplets may contribute significantly to the aerosol direct effect.^{8,11,13} Our previous study did not find evidence of drying-induced BrC formation in ambient particles,¹⁴ most likely due to the low concentration of dicarbonyls in ambient particles relative to laboratory studies that have observed such a phenomenon.²⁵ However, in clouds, the partitioning of BrC precursors is expected to be several orders of magnitude higher, due to the high LWC.^{59,60} For example, Gly + MeGly can constitute up to $\sim 10\%$ of WSOC in clouds and fogs,^{2,61} demonstrating the greater potential for drying-induced BrC formation compared to aqueous particles. Fig. 4b shows the ΔAbs value ($\Delta Abs = Abs_{dry} - Abs_{byp}$) for all five cloud water samples. Similar to our prior results for the eastern US aerosol, we did not observe evidence of BrC formation in the aerosolized cloud water samples as a result of drying within our method detection limit of 1×10^{-3} A.U.¹⁴ There are a few possible explanations for this observation. As mentioned earlier, airmasses reaching WFM were likely aged, suggesting that any secondary BrC production may have occurred during transport to WFM, largely depleting the airmass of BrC precursors.^{18,25,62} Photobleaching (decreasing absorbance) of secondary BrC



occurs on time scales of minutes to days,^{63–65} so any BrC formation upwind of WFM would likely have been affected by this process.^{65,66} Another possible explanation for the lack of BrC formation in our experiments is that the drying RH of 50% may not be sufficient to produce substantial BrC. In a previous study, we observed BrC formation in aqueous glyoxal/(NH₄)₂SO₄ particles when dried to ~30% RH, but not when the particles were dried to ~50% RH.¹⁴ In the same study, we observed BrC formation from MeGly/(NH₄)₂SO₄ when aqueous particles were dried to both 50% and 30% RH, but a significant enhancement at the lower RH level. A drying time of ~7 s was sufficient to produce BrC formation in aqueous particles undergoing drying in previous laboratory studies.^{11,14} Notably, in our previous study conducted with a nearly identical experimental setup, we observed BrC formation in aqueous glyoxal and methylglyoxal droplets with only ~7 s of drying.¹⁴ However, the effect of variable drying times on BrC formation has not been systematically studied.

3.2 Model results

Simulations from GECKO-A were used to predict the reversible partitioning of VOC oxidation products to cloud water after the equivalent of two days of gas-phase photooxidation. There is a wide distribution of WSOC_{dry}/WSOC_{cw} predicted for the VOC oxidation products, with precursor molecule identity and oxidation regime (*i.e.*, VOC/NO_x) both contributing to this variability (Fig. 5). While the simulations are simplistic in that they do not account for reactions of the organics in CW, which can shift the partitioning of compounds produced or consumed in the aqueous phase, they are instructive in a number of ways.

For most of the simulated VOCs, photooxidation under high-NO_x conditions leads to a decrease in the predicted WSOC_{dry}/WSOC_{cw} due to an increase in the relative abundance of volatile oxidation products (compounds like Gly and MeGly). This is most evident in the single ring aromatic simulations, which all show <2% of the WSOC that partitions to CW remains in the condensed phase after drying (mole fraction basis). The simulations also indicate far greater production of higher volatility products from these compounds compared to the species with much higher predicted WSOC_{dry}/WSOC_{cw}, for example, C₂₂- and C₁₈-alkanes.

The simulations are interesting to contrast with the lab measurements of WSOC_{dry}/WSOC_{cw} in the WFM samples (green diamonds in Fig. 5). The measured WSOC_{dry}/WSOC_{cw} shows a relatively narrow distribution, which is perhaps not surprising given that the CW samples were collected over such a short period of time (12-Aug-2017 to 18-Aug-2017, Table S1†). While we observed systematic evaporation of some WSOC in all five CW samples (11% on average), the majority of the WSOC compounds had sufficiently low volatility to remain in the condensed phase upon drying.

Broader analysis of WFM data suggests a wider distribution of volatilities in the CW than our five samples showed (blue crosses in Fig. 5). The data presented are summertime monthly averages (June–September, 2009–2018) of the ratio of organic aerosol concentrations (OA_{AER}) measured downwind of the WFM summit relative to CW organics (both concentrations expressed per m³ of air for direct comparison).⁶⁷ While the summit and downwind sites are not always sampling the same airmass, simultaneous O₃ measurements suggest that the prevailing conditions at the summit where CW collection occurs and the aerosol sampling site are representative (Fig. S3, ESI†). The monthly average OA_{AER}/OA_{CW} ratio (blue crosses in Fig. 5) has a wide range of values, from a low of 0.3 to a high of ~1, with an average value of 0.6. This reflects influence from the oxidation of numerous VOCs, wide-ranging atmospheric conditions, and variable aging times during transport to WFM. Even when the site conducting aerosol sampling is downwind of the summit, there can be difficulties directly connecting the CW and the aerosol composition measurements due to droplet deposition.^{15,20} Thus, the values of the blue crosses in Fig. 5 have a high level of uncertainty and would be biased low if deposition removed organics during transport from the summit to the downwind site. Nevertheless, the lower OA concentrations in nearly all months suggests the evaporation of organics from clouds at this site is a systematic occurrence during the summertime. The modeling and long-term WFM results in Fig. 5 suggest that a much broader range of WSOC_{dry}/WSOC_{cw} values would be observed if the laboratory measurements were expanded to include cloud water samples influenced by a broader range of source regions and atmospheric ageing conditions.

These results reveal the importance of clouds in the uptake of oxidized VOCs, which demonstrates significant potential to produce aqSOA. A lower predicted WSOC_{dry}/WSOC_{cw} indicates a more volatile product distribution and thus a greater potential to form aqSOA in cloud water. Note

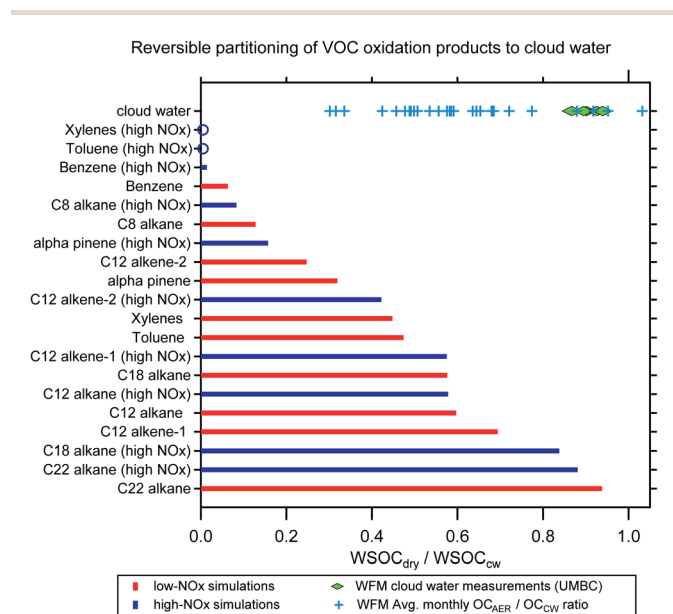


Fig. 5 GECKO model predictions of WSOC_{dry}/WSOC_{cw} for the VOC photooxidation products (red and blue bars). The model results were compared with the laboratory cloud water measurements (green diamonds) and with the ratio of organic aerosol to cloud water organics measured at WFM (summertime monthly averages from 2009–2018, blue crosses).



that the GECKO modeling results include vastly more oxygenated organics than have been identified through molecular measurements, in agreement with the finding that the majority of WSOC in cloud water is unidentified on a molecular level.² Together, this suggests that aqSOA formation may be even more substantial than estimates using known precursors.

4. Conclusions

Cloud cycling is an important process that leads to physical and chemical transformations of dissolved organic carbon. We found the drying (to ~50% RH) of cloud water droplets resulted in ~11% evaporative loss of WSOC. This observation confirms that organics in cloud water are reversibly partitioned to the aqueous phase, and likely represents a lower bound on the evaporation of organics during an actual cloud cycle (due to experimental measurement constraints). The relatively low evaporative loss of WSOC (~11%) compared to the GECKO simulations also suggests the occurrence of aqueous processing of organics in cloud water during transport to WFM. Formic and acetic acid account for a small fraction, 19% on average, of this reversibly partitioned organic material. This may be an important process that affects the distribution and lifetime of reactive carbon in the atmosphere. The lifetime of semi-volatile organics is lower in the gas phase than in the condensed phase, so reversible partitioning to cloud water may reduce deposition losses and enhance the transport of these compounds.⁵³ Although our laboratory analysis included a relatively small sample size ($n = 5$), the results from long-term observations at WFM show systematically lower organic aerosol concentrations downwind of the summit than in cloud water. This supports our laboratory measurements and suggests the nearly constant presence of volatile WSOC reversibly partitioned to cloud water. GECKO box model simulations predict abundant formation of compounds that reversibly partition to cloud water from the photooxidation of ten different VOCs under high- and low-NO_x conditions. However, the relatively low evaporative loss of WSOC (~11%) compared to the GECKO simulations also suggests the occurrence of aqueous processing in cloud water during transport to WFM.

No evidence of drying-induced BrC formation was observed in the cloud water samples. The most likely reason appears to be the air mass reaching WFM may have been depleted of BrC precursors.^{18,25} Many studies have reported the loss of absorbance of BrC due to photooxidative aging, which could also contribute to the depletion of BrC and BrC precursors during transport.⁶² Studies have confirmed the need to account for photobleaching of BrC in order to accurately constrain BrC and global radiative balance, so this may have had an impact on the observations.⁶⁶

The work highlights the potential importance of the large unidentified fraction of WSOC in cloud and fog water, as well, suggesting that current detailed molecular measurements of cloud water will likely be incomplete and may not capture broader trends in the behavior of organics.

Conflicts of interest

The authors declare no competing financial interest.

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