Reversible and irreversible gas-particle partitioning of dicarbonyl compounds observed in the real atmosphere

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Abstract. Glyoxal and methylglyoxal are vital carbonyl compounds in the atmosphere and play substantial roles in radical cycling and ozone formation. The partitioning process of glyoxal and methylglyoxal between the gas and particle phase via reversible and irreversible pathways could efficiently contribute to secondary organic aerosol (SOA) formation. However, the relative importance of two partitioning pathways still remain elusive, especially in the real atmosphere. In this study, we launched five field observations in different seasons and simultaneously measured glyoxal and methylglyoxal in the gas and particle phase. The field-measured gas-particle partitioning coefficients were 5–7 magnitudes higher than the theoretical ones, indicating the significant roles of reversible and irreversible pathways in the partitioning process. The particulate concentration of dicarboxyls and product distribution via the two pathways were further investigated using a box model coupled with the corresponding kinetic mechanisms. We recommended the irreversible reactive uptake coefficient $\gamma$ for glyoxal and methylglyoxal in different seasons in the real atmosphere, and the average value of $8.0 \times 10^{-3}$ for glyoxal and $2.0 \times 10^{-3}$ for methylglyoxal best represented the the loss of gaseous dicarboxyls by irreversible gas-particle partitioning processes. Compared to the reversible pathways, the irreversible pathways played a dominant role, with a proportion of more than 90% in the gas-particle partitioning process in the real atmosphere and the proportion was significantly influenced by relative humidity and inorganic components in aerosols. However, the reversible pathways were also substantial, especially in winter, with a proportion of more than 10%. These two pathways of dicarboxyls jointly contributed to more than 25% of SOAs in the real atmosphere. To our knowledge, this study was the first to systemically examine both reversible and irreversible pathways in the ambient atmosphere, strove to narrow the gap between model simulations and field-measured gas-particle partitioning coefficients, and revealed the importance of gas-particle processes for dicarboxyls in SOA formation.

1 Introduction

Glyoxal and methylglyoxal, the simplest $\alpha$-dicarbonyls, are recognized as being of great importance in atmospheric chemistry due to their unique physicochemical properties. The $\alpha$-dicarbonyl functionality increases their water solubility and reactivity more than expected. The traditional opinion is that methylglyoxal is less reactive compared to glyoxal due to its unreactive
methyl substitution, while a very recent study noted that methylglyoxal could be more reactive under an atmospheric-relevant concentration (Li et al., 2021). Overall, both of them play crucial roles in radiation balance, air quality, brown carbon formation, and SOA formation (Laskin et al., 2015; Qiu et al., 2020). Moreover, as major carcinogenic and genotoxic compounds, dicarbonyls can cause serious damage to human health. They have relatively limited primary sources, except for biomass burning and biofuel combustion (Zarzana et al., 2018; Zarzana et al., 2017), compared to secondary formation that occurs with photooxidation of both biogenic volatile organic compounds (VOCs), such as isoprene, and anthropogenic VOCs, such as aromatic hydrocarbons (Lv et al., 2019). Considering the atmospheric sink, glyoxal and methylglyoxal can be lost in the gas phase by photochemical reactions, oxidation by OH radicals, and dry deposition; however, there is still a missing sink for the two dicarbonyls (Volkamer et al., 2007).

Gas-particle partitioning was recently found to be the most important removal pathway for both glyoxal and methylglyoxal, especially in regions like Beijing with high particulate matter (PM) pollution that provides sufficient aerosol surface area. Although with relatively high vapor pressure, glyoxal and methylglyoxal can efficiently partition into the particle phase due to their α-dicarbonyl functionality. Upon physical adsorption to the particle phase, they can undergo various chemical reactions and subsequently form larger-molecular-weight products retained in the condensed phase. Moreover, chemical reactions can in turn accelerate the physical adsorption and greatly contribute to the formation and growth of atmospheric particulate matter. The chemical reactions occurring in the gas-particle partitioning processes can be divided into reversible pathways, including reversible hydration and self-oligomerization, and irreversible pathways, which can be driven by oxidative radicals and other reactive compounds. These processes can also efficiently explain observed aerosols properties – including relatively high oxygenation levels, compositions like organic acids and oligomers, and higher light absorption – that cannot be explained by traditional absorptive models of gas-particle partitioning (Pankow, 1994; Pankow and James, 1994; Odum et al., 1996).

Many laboratory and model studies have made a great effort to investigate the reversible and irreversible pathways of dicarbonyls to further understand their gas-particle partitioning mechanisms and reveal their contribution to SOA formation. Fu et al. (2008) found that the modeled SOA concentrations were largely increased when accounting for irreversible uptake of dicarbonyls in the GEOS-Chem model. Considering the surface-controlled reactive uptake of dicarbonyls into the CMAQ model, the aerosol uptake of dicarbonyls accounted for more than 45% of total SOA in the eastern US (Ying et al., 2015); similarly, the contribution of glyoxal and methylglyoxal to SOA formation in China was 14% to 25% and 23% to 28%, respectively (Hu et al., 2017). Although reversible and irreversible pathways of dicarbonyls have been separately investigated in previous studies, solely incorporating just one pathway into models could lead to a large discrepancy between model results and observational data, highlighting the importance of comprehensively considering both reversible and irreversible pathways when quantifying the gas-particle partitioning process of dicarbonyls (Li et al., 2014; Hu et al., 2017; Ling et al., 2020). Despite increasing interest in dicarbonyls and their gas-particle partitioning processes, the detailed chemical mechanisms of
two partitioning pathways remain poorly understood. First, previous studies have exposed seed particles to high concentration levels of dicarbonyl vapors, from hundred ppb to ppm levels, or used bulk samples; thus, their applicability to the real atmosphere requires further validation. Second, prior studies always used one constant coefficient to present all heterogeneous processes occurring on the aerosol, which neglects the influencing factors in real atmospheric partitioning processes. Further studies have shown that the two pathways in the gas-particle partitioning process for glyoxal and methylglyoxal are rather complex, and their relative contribution to the partitioning process can be influenced by many factors such as relative humidity (Curry et al., 2018; Shen et al., 2018), particle acidity (Liggio et al., 2005b; Shi et al., 2020), and particle organic/inorganic components (Kampf et al., 2013). However, there persist controversies in the specific partitioning mechanisms of glyoxal and methylglyoxal, especially conflicting views on their role in SOA formation, which urgently warrants further investigation.

In this study, five field observations were launched over urban Beijing in four seasons, and glyoxal and methylglyoxal in the gas and particle phase were simultaneously measured. Beijing, as the political center of China, is the most prosperous city with numerous key environmental issues. Chen et al. (2021) found that the average concentration of dicarbonyls in Beijing is lowest among key regions with relatively higher PM$_{2.5}$ concentrations, indicating there is a more efficient partitioning process of dicarbonyls. Thus, it is more environmentally significant to discuss the gas-particle partitioning processes in urban Beijing. These processes are divided into two pathways: reversible and irreversible. On the basis of field-measured data, we could estimate the product distribution, main influencing factors, and relative importance of the two gas-particle partitioning pathways for glyoxal and methylglyoxal in the real atmosphere.

2 MATERIALS AND METHOD

2.1 Field sampling and analysis

We performed field observations on the roof of a six-story teaching building (26 m above the ground) on the Peking University campus (39.992°N, 116.304°E) in northwest urban Beijing. The field observations in this study were launched during four different seasons from 2019 to 2021.

Gaseous carbonyls were collected by adsorption reactions in a 2,4-dinitrophenyl hydrazine (DNPH) cartridge (Sep-Pak; Waters Corporation). The air samples were first passed through an ozone scrubber (Sep-Pak; Waters Corporation) to eliminate interference by ozone and then trapped in the DNPH cartridge. To prevent deliquescence of the potassium iodide in the ozone scrubber, the air samples were mixed with ultrapure nitrogen before pumped into the sampling tubing.

Particulate carbonyls were collected by a four-channel ambient particles sampler (TH-16A, Wuhan Tianhong) with Teflon filters and quartz filters (47 mm, Whatman). The Teflon filters were used to measure the mass concentration of collected PM$_{2.5}$ and the quartz filters were used for carbonyl analysis. Detailed information about field sampling and analysis were provided.
in our previous studies (Rao et al., 2016; Qian et al., 2019).

The meteorological station was co-located at our sampling site and provided meteorological parameters. Common trace gases like NO/NO₂, SO₂, CO, and O₃ were detected online by Thermo 42i, 43i, 48i, and 49i analyzers, respectively. A TEOM 1400A analyzer was applied to measure the mass concentrations of PM_{2.5} and PM_{10}, the results of which were consistent with the PM_{2.5} weighing results (Fig. S1). The time solution for all of the above data was 1 min. Detailed information about these five observations is shown in Table S1.

2.2 Sample extraction and analysis

The gaseous carbonyl samples were eluted with acetonitrile (HPLC/GC-MS grade), and the particulate carbonyl samples on quartz filter were eluted with acidic DNPH solutions in the flask and then were shaken for 3 h at 4 °C with a rotation rate of 180 rpm in an oscillator (Shanghai Zhicheng ZYW 103D). The derived solutions were placed in darkness for 12-24 h to ensure complete derivatization, and then they were analyzed by high-performance liquid chromatography-ultraviolet (HPLC-UV) for separation and detection. The limit of detection (LOD) of this method was 50 pptv for gaseous carbonyls and 1 ng·m⁻³ for particulate carbonyls. They were calibrated using a mixing standard solution with a concentration range of 0.1–10 µM, and the linearity was indicated by a correlation of determination (r²) of at least 0.999. The detailed analysis method was presented in our previous study (Wang et al., 2009).

The Teflon samples were also extracted by deionized water using an ultrasonic bath for 30 min at room temperature. The extracted solutions were analyzed by ion chromatography (Integrion and Dionex ICS 2000, USA) to measure the water-soluble inorganic compounds (Na⁺, NH₄⁺, K⁺, Mg²⁺, Ca²⁺, Cl⁻, NO₃⁻, and SO₄²⁻) and low-molecular-weight organic acids (formate, acetate, and oxalate) in aerosols.

2.3 Estimation of effective partitioning coefficient

To estimate the effective partitioning process of gas-phase carbonyls to the particle phase, we could use Pankow’s absorptive partitioning theory for the gas-organic phase (Eqs. (1), (2)) (Odum et al., 1996) and Henry’s law for the gas-liquid phase (Eq. (3)):

\[ K_p^{eq} = \frac{C_p}{C_{eq-TSP}} \] (1)

\[ K_{eq} = \frac{RT_{cm}}{10^{M/2.4C_p}} \] (2)

\[ K_H = 10^\frac{C_p}{\rho_p \cdot M \cdot ALWC \cdot C_{water}} \] (3)

In Eq. (1), \( K_p^{eq} \) (m³·µg⁻¹) is the field-measured gas-particle partitioning coefficient; \( C_p \) (µg·m⁻³) and \( C_{eq} \) (µg·m⁻³) are the concentrations of dicarbonyls in the particle and gas phase, respectively; and TSP (µg·m⁻³) is the mass concentration of
suspended particles (mass concentrations of PM$_{2.5}$ were used in this study). In Eq. (2), $K_{p}$ ($m^3\cdot \mu g^{-1}$) are the theoretical gas-particle partitioning coefficients determined by Pankow’s absorptive model, $f_{om}$ is the absorbing fraction of total particulate mass, $MW_{OM}$ (g·mol$^{-1}$) is the mean molecular weight of the organic phase, and $\zeta$ is the activity coefficient of target compounds.

In the estimation of $K_{p}$ in this study, $f_{om}$ and $\zeta$ are unity and $MW_{OM} = 200$ g·mol$^{-1}$, as used in previous studies (Healy et al., 2008; Williams et al., 2010; Xie et al., 2014; Shen et al., 2018), and $p^{0}_1$(Pa) is the supercooled vapor pressure of compounds as a pure liquid at temperature $T$, which is calculated by the extended aerosol inorganic model (E-AIM, http://www.aim.env.uea.ac.uk/aim/ddbst/pcalc_main.php) (Clegg et al., 1998). In Eq. (3), $K_{H}$ (M·atm$^{-1}$) is the field-derived effective Henry’s law coefficient; $c_{p}$ (μg·m$^{-3}$) and $c_{g}$ (atm) are particle- and gas-phase concentrations of carbonyls, respectively; ALWC (μg·m$^{-3}$) is the aerosol liquid water content calculated by the thermodynamic model ISORROPIA-Ⅱ (forward model, metastable state), the results of which are comparable to the actual measured contents confirmed by previous studies (Guo et al., 2015).

3 RESULT AND DISCUSSION

3.1 Observation results and partitioning coefficients calculation

3.1.1 Dicarbonyls in the gas and particle phase

We launched five field observations in different seasons. Table S1 details the information about the field observations, including observation periods, sample volume, and meteorological parameters. We totally collected 387 gas-phase samples and 130 particle-phase samples in four seasons. In these samples, carbonyls were simultaneously measured in both gas phase and particle phase. Ten carbonyls were measured in the gas phase and six carbonyls were measured in the particle phase. In this study, we mainly discuss the gas-particle partitioning processes of glyoxal and methylglyoxal because of their significant roles in atmospheric chemistry.

Figure 1 and Table 1 show the temporal characteristics and seasonal variation of glyoxal and methylglyoxal, respectively. Gaseous dicarbonyls showed obvious seasonal variation. Concentrations in summer ($0.99 \pm 0.59$ ppbv) were generally much higher than in other seasons, followed by autumn and spring, and the concentrations in winter were the lowest. This seasonal variation could be partly attributed to the higher temperature and more intensive radiation in summer, which could greatly enhance the secondary formation of gaseous carbonyls via photochemical reactions. The diurnal variation in the dicarbonyls during summer support this interpretation of the data; gas-phase dicarbonyls exhibited obviously diurnal variations in summer, whereas this variation was irregular in other seasons (Fig. S2). The concentration levels of gaseous dicarbonyl in summer rapidly increased after sunrise, remained relatively high during the daytime (12:00–14:00), and then decreased at dusk.

Although methylglyoxal has a shorter lifetime compare to glyoxal (GL 2.9 h vs. MG 1.6 h) (Fu et al., 2008), its gas-phase
concentration levels were generally higher than those of glyoxal, consistent with previous studies (Rao et al., 2016; Mitsuishi et al., 2018; Qian et al., 2019), mainly due to the relatively larger production from isoprene and acetone for methylglyoxal. The concentrations of particulate dicarbonyls were an order of magnitude smaller than the gaseous concentrations using the unit of nanogram per cubic meter of air (ng/m$^3$ air). The average particulate glyoxal and methylglyoxal were 19.37 and 11.24 ng/m$^3$, respectively, which were slightly higher than previously reported values (Zhu et al., 2018; Shen et al., 2018; Cui et al., 2021; Qian et al., 2019). Dicarbonyls measured in the particle phase also showed obvious seasonal variation. The particulate concentrations of the two dicarbonyls in winter (43.38 ± 32.42 ng/m$^3$ air) were 2–2.3 times higher than those in other seasons, suggesting that the dicarbonyls were more favored into the particle phase in winter. Moreover, particulate dicarbonyls in different seasons exhibited the same diurnal variation (Fig. S2). The particulate concentrations of dicarbonyls in daytime were generally higher than those in nighttime, especially in winter.

### 3.1.2 Gas-particle partitioning coefficient

Dicarbonyls could partition between gas and aerosol phases or the liquid phase, following Pankow’s absorptive partitioning theory or Henry’s law, respectively, as listed in Table 2. Both gas-particle partitioning coefficient ($K_p^f$) and effective Henry’s law coefficient ($K_H^f$) were calculated on the basis of field-measured data and were in the range of $10^{-4}$–$10^{-2}$ m$^3$·µg$^{-1}$ and $10^6$–$10^9$·M·atm$^{-1}$, respectively. The partitioning coefficient values of the two dicarbonyls exhibited the same seasonal variation, as winter and spring > autumn > summer. A higher aerosol concentration accompanied by higher aerosol surface area concentration and lower relative humidity resulted in a higher partitioning coefficient in winter and spring, when heavy pollution and sandstorms always occurred. In the case of temperature variation varied from 265.53 K to 310.75 K in different seasons, lower temperature promoted the partitioning processes as $K_p^f$ values for the dicarbonyls and temperature showed negative correlation with significant difference ($p < 0.001$) (Fig. S3). Moreover, The $K_p^f$ and $K_H^f$ values of glyoxal were always higher than those of methylglyoxal, implying the former was more likely to partition to the particle phase; this could be attributed to their different structures. Glyoxal were more soluble and reactive because of the adjacent electron-poor aldehydic carbons, whereas methylglyoxal was more stable due to the reduced electron-deficient ketone moiety (Kroll et al., 2005).

Both $K_p^f$ and $K_H^f$ were relatively closed to those found in previous field-measured studies (Shen et al., 2018; Qian et al., 2019; Cui et al., 2021). However, compared with the theoretical partitioning coefficients $K_p^t$ calculated by Pankow’s absorptive theory, $K_p^f$ values were approximately 5–7 orders of magnitudes higher than the corresponding $K_p^t$ values. The influencing factors in Pankow’s absorptive model, like the activity coefficient $\zeta$ or absorbing fraction $f_{\text{am}}$, could not explain this great difference between the field-measured values and the theoretical ones. The underestimation of gas-particle partitioning coefficients can be attributed to the misidentification of condensed phase species produced in heterogeneous
chemical reactions. Similarly, $K_H^f$ values were approximately 2–5 orders of magnitudes higher than the theoretical Henry's law coefficient $K_H^l$ calculated in pure water. The discrepancy could be explained by the complex components in aerosol liquid water. Figure S4 presents the Sefcikenow plot of dicarbonyls versus aqueous sulfate, nitrate, and ammonia (SNA) concentration in aerosol. The negative salting constant indicated the “salting in” effects, which could result in exponential solubility, for both glyoxal and methylglyoxal in the real atmospheric. Moreover, both $K_H^f$ and $K_H^l$ of dicarbonyls were more than one magnitude higher than the reported laboratory partitioning coefficient values from chamber experiments (Healy et al., 2008; Healy et al., 2009), indicating that the real atmosphere is more favorable for the partitioning of gaseous dicarbonyls to the particle phase. Actual atmospheric environment conditions and complex particle compositions, such as higher ionic strength, could greatly affect the partitioning process and the chemical reactions in the aerosols.

To narrow the large discrepancy between field-measured partitioning coefficients and theoretical (or laboratory) ones, we needed to further investigate the mechanism and product distribution of chemical reactions occurring in the aerosols during the partitioning processes. The products of the reversible pathway mostly have lower saturated vapor pressure, and thus leading to higher partitioning coefficients compared to monomer dicarbonyls. For example, the calculated vapor pressures of the products of glyoxal hydration and dimerization are, respectively, 5 and 10 orders of magnitudes less than that of glyoxal monomer (Hastings et al., 2005). Moreover, the products of the irreversible pathway, such as organic acids produced in radical chemistry, also have lower vapor pressure and efficiently contribute to the underestimation of partitioning coefficients. The following sections further discuss the mechanism and product distribution of reversible and irreversible pathways to explain the partitioning process of dicarbonyls.

### 3.2 Reversible pathways

Gas-particle partitioning of dicarbonyls via a reversible pathway mainly consists of hydration and self-oligomerization. Since glyoxal and methylglyoxal had high water solubility and reactivity, they could easily dissolve into aerosol liquid water and form hydrates, which are more reactive than their counterparts and could participate in continuous reactions to form higher-molecular-weight oligomers. Hemiacetal/acetal formation (Loeffler et al., 2006) and aldol condensation (Haan et al., 2009) are the most thermodynamically favored oligomer reactions for glyoxal and methylglyoxal, respectively. The proposed mechanism for the reversible formation of glyoxal and methylglyoxal in aerosols is shown in Fig. S5. Overall, since the products of the reversible pathway, including hydrates and oligomers, are thermodynamically unstable and could easily revert to their original monomer form during extraction and analysis, their total concentration could be presented as the measured particle-phase dicarbonyls from sampled quartz filters. Since glyoxal and methylglyoxal have similar trend under different conditions, we focused on the total concentration of the two dicarbonyls in the following discussion. As shown in Fig. 2a, the particulate concentration of dicarbonyls via a reversible pathway is strongly dependent on RH. It increased significantly when
RH increased from <10% to 60%; however, from 60% to 80% RH, it exhibited the opposite trend and decreased with increasing RH. Moreover, under high RH conditions, the particulate concentration of dicarbonyls via a reversible pathway had a strong and positive dependence on particle acidity (pH). The product distribution of the reversible formation could well explain this phenomenon.

To roughly estimate the product distribution of the reversible pathway in the real atmosphere, we simplified reaction mechanisms and calculated the product distribution on the basis of the equilibrium constant reported in previous literature (Table S2). Generally, more dicarbonyls existed in oligomer forms (83.5% for glyoxal and 80.8% for methylglyoxal) than in hydrate forms (16.3% for glyoxal and 20.8% for methylglyoxal) in the reversible formation. Moreover, their distribution exhibited obvious seasonal variations. Summer had the highest proportion of hydrate forms (52.80% for GL and 35.10% for MG), while winter had the highest proportion of oligomer forms (86.15% for GL and 86.31% for MG). Detailed information is shown in Table S3. The seasonal variation could be attributed to the RH in different seasons – relatively high in summer and low in winter. As shown in Fig. 2b, the product distribution of the reversible formation has a strong dependence on RH. The proportion of dicarbonyls in hydrate forms increased with increasing RH and could reach more than 75% in high RH, while the proportion of dicarbonyls in oligomer forms exhibited the opposite trend. Hydrates play a dominant role in dilute solutions under high RH conditions with a relatively high aerosol liquid water concentration, which might hinder oligomer formation. And large quantities of oligomers, including dimers and trimers, would form until the aerosol liquid concentration became greater than 1 M (Liggio et al., 2005b) when RH decreased.

Combined with the vapor pressure of dominant products, their gas-particle partitioning coefficient can be roughly estimated and can effectively fit the field-measured values, as shown in Fig. 2c. The estimated gas-particle partitioning coefficients in this study are five orders of magnitude higher than the theoretical ones but still 1–2 orders of magnitude lower than the field-measured coefficients, especially in winter. The difference between the estimated partitioning coefficients and the field-measured ones suggests that the current understanding of the equilibrium in reversible formations cannot reasonably explain the gas-particle partitioning processes of dicarbonyls. There still exist extra pathways of reversible formation. Cross-oligomerization of glyoxal and methylglyoxal is nonnegligible and could form similar molecular structure products and contribute to SOA yield (Schwier et al., 2010). Esterification and amination of diols also occur in aerosol liquid water but are negligible compared to hydration and polymerization (Zhao et al., 2006). However, these reactions are not further discussed here. The hydrates and oligomers mentioned above are the dominant forms of glyoxal/methylglyoxal in the particle phase, while the higher molecular oligomers up to nonamer could also exist with a relatively smaller but still significant fraction at equilibrium. Although the reactions are thermodynamically reversible, upon evaporation of the aerosol liquid water, the oligomer formation is faster than the evaporation of dehydrated dicarbonyls, and the dicarbonyl evaporation is limited (Liggio et al., 2005b; Loeffler et al., 2006). This results in relatively stable oligomers and yielding SOA. Moreover, other nucleophilic
species may also form oligomers with glyoxal and methylglyoxal and effectively prevent their evaporation. Besides reversible pathways, higher carbon number products with lower volatility were mainly formed through irreversible pathways, like radical reactions (e.g., OH radicals), which are fully discussed in the next section.

3.3 Irreversible pathways

3.3.1 Irreversible pathways driven by hydroxyl radicals

Reactive uptake driven by hydroxyl radicals (OH) is the dominant process for glyoxal and methylglyoxal in their irreversible gas-particle partitioning pathways. Compared to other irreversible pathways, like imidazole formation, glyoxal/methylglyoxal + OH chemistry occurs on much shorter timescales (Teich et al., 2016). The reaction is the initial step for most radical-based chemistry of glyoxal/methylglyoxal and has been proven to be an important source of SOA in both cloud/fog droplets and wet aerosols (Tan et al., 2012; Lim et al., 2013), producing low-volatility products such as organic acids, large multifunctional humic-like substances, and oligomers. The proposed mechanism for the irreversible pathway of glyoxal and methylglyoxal driven by hydroxyl radicals in aerosols is shown in Fig. S6. The OH radicals in aerosol liquid water are mainly from the direct uptake of gas-phase OH radicals with a Henry’s law constant of 30 M/atm (Faust and Allen, 1993) and Fenton reactions, and Fenton reactions are closely related to hydrogen peroxide, iron ions, and manganese ions in the particle phase. The sources of OH radicals are one of the major uncertainties in SOA formation (Ervens et al., 2014).

The irreversible reactive uptake coefficient $\gamma$ could efficiently describe the irreversible pathway of the gas-particle partitioning process of dicarbonyls driven by OH radicals. We could estimate the reactive uptake coefficients $\gamma$ based on the effective Henry’s constant via theory calculation (Hanson et al., 1994; Curry et al., 2018) and then calculate the effective uptake rate $k_{eff,\text{uptake}}$ following Eqs. (4)–(7):

$$\frac{1}{\gamma} = \frac{1}{a} + \frac{\gamma}{4RT^{1/2}D_{aq}^{1/2}\coth(q+1/4)}$$  \hspace{1cm} (4)

$$v = \frac{8RT}{\pi M_X}$$  \hspace{1cm} (5)

$$q=R_p/l=R_p/\sqrt[3]{4M_X/\pi}$$  \hspace{1cm} (6)

$$k_{eff,\text{uptake}} = v \times \gamma \times A_{surf}$$  \hspace{1cm} (7)

where $\gamma$ is the dimensionless uptake coefficient, $v$ (m s$^{-1}$) is the gas-phase thermal velocity of glyoxal/methylglyoxal, $D_{aq}$ (m$^2$ s$^{-1}$) is the diffusion coefficient in the liquid phase, $a$ is the dimensionless mass accommodation coefficient, $H^*$ (M atm$^{-1}$) is the effective Henry’s law constant calculated by field-measured data in Table 2, $R$ is the universal gas constant, $k^f$ (s$^{-1}$) is the first-order aqueous loss rate, $M_X$ (kg mol$^{-1}$) is the average molar mass of gas-phase dicarbonyls, $q$ is the parameter for measuring in-particle diffusion limitations, $R_p$ (m) is the particle radius, $l$ (m) is the diffusion reactive length, $k_{eff,\text{uptake}}$ (s$^{-1}$) is
the effective uptake rate, and $A_{afl}(m^2 \cdot m^{-3})$ is the aerosol surface area density. This formulation is based on the effective Henry’s law constant under high RH conditions (RH > 40%). Moreover, the formulation describes the reactive uptake due to irreversible multiple-phase loss processes in the presence of OH. The uncertainty in the $\gamma$ calculation is mainly attributed to the uncertainty in OH concentration, which was $3 \times 10^{-12}$ M on average and varied from $5.5 \times 10^{-14}$ to $8 \times 10^{-12}$ M (Herrmann et al., 2010).

The calculated $\gamma$ and $k_{eff,uptake}$ values for different seasons are listed in Table 3. The reactive uptake coefficients of glyoxal were in the range $10^{-4}$–$10^{-2}$, and the average value of $8.0 \times 10^{-3}$ in this study was closed to the ones representing the loss of glyoxal by surface uptake during the KORUS-AQ campaign in a very recent studies (Kim et al., 2022). And the value slightly exceeded the one commonly used in model simulations ($\gamma = 2.9 \times 10^{-3}$), which was based on an experimental study for (NH$_4$)$_2$SO$_4$ aerosols at 55% RH (Liggio et al., 2005a), and also far outweighs the uptake coefficients of glyoxal on clean and acidic gas-aged mineral particles ($\gamma = 10^{-4}$–$10^{-5}$) (Shen et al., 2016), implying that a real atmospheric aerosol provides a far more reactive interface for physiochemical processes than that of mineral particles. Moreover, the reactive uptake coefficients of methylglyoxal were slightly lower than those for glyoxal, with an average value of $2.0 \times 10^{-3}$. Conflicting with previous experimental results, methylglyoxal exhibited unexpected salting-in effects in real atmospheric particles and had much higher uptake coefficients, which could be attributed to the increased reactivity of methylglyoxal with a high uptake coefficient under an atmospheric relevant concentration (Li et al., 2021). The $\gamma$ values for both glyoxal and methylglyoxal exhibited similar seasonal variations, which were lowest in summer and reached their highest in winter. This seasonal variation could be attributed to RH variation and particle composition. Moreover, the effective uptake rate ($k_{eff,uptake}$), which is regarded as a pseudo-first-order reaction rate, is a net result of competition between reversible and irreversible processes, and it varied from $10^{-4}$ s$^{-1}$ to $10^{5}$ s$^{-1}$ in the real atmosphere in this study. As shown in Fig. 3a, the negative dependence of $k_{eff,uptake}$ on RH also confirmed that the irreversible uptake of dicarbonyls could be inhibited in high RH conditions. And the relatively low SNA concentration under high RH conditions also attenuated the irreversible uptake as the weakening of ion effects (Figure 3b).

Whereas, for a given RH, uptake coefficients $\gamma$ for both glyoxal and methylglyoxal showed a weak dependence on the ratio of SNA (S:A and S:N) with significant scatter (Fig. S7).

Moreover, it was worth noting that under extremely low RH (<40%), the aerosol was not completely deliquescent, and the uptake coefficients based on Henry’s law could not explain the irreversible pathways. Previous research indicated that the irreversible uptake of dicarbonyls could still occur under a low RH condition (Liggio et al., 2005a; De Haan et al., 2018), and that these uptake values were generally lower due to the inefficient reactive uptake process onto the crystallized aerosols.

### 3.3.2 Reactive uptake of dicarbonyl compounds

We could not directly measure the particulate concentration of dicarbonyls via an irreversible pathway, as the dicarbonyls irreversibly reacted with oxidative radicals on aerosols. To quantitively evaluate the contribution of the irreversible pathway...
of dicarbonyls, we calculated their average concentration based on Eqs. (S1)–(S5) in the Supplement with the calculated $\gamma$ values in this study. The samples estimated here were collected under high RH conditions (RH > 40%) because of the calculation limitation of irreversible uptake coefficients.

The total particulate concentration of glyoxal and methylglyoxal via irreversible pathway varied from several to more than 100 nanograms per microgram PM$_{2.5}$ (ng/μg PM$_{2.5}$), and it was strongly dependent on RH, as shown in Fig. 3c. Since the irreversible uptake coefficients $\gamma$ of dicarbonyls tended to decrease with increasing RH due to the “salting-in” effects, both glyoxal and methylglyoxal in this study exhibited high solubility in low RH conditions, where the aerosol water was coupled with concentrated inorganic solutions and relatively high ionic strength. Moreover, the total particulate concentrations of dicarbonyls were positively dependent on particle acidity under high RH conditions, while there were no obvious correlations under low RH conditions.

To further discuss the product distribution of the reaction of glyoxal/methylglyoxal with hydroxyl radicals, we used the kinetic mechanisms of glyoxal/methylglyoxal + OH chemistry proposed by Lim et al. (2013) on the basis of a 0-D box model with a steady-state approach. The average OH radical concentration setting in the modeling was $3.2 \times 10^{-12}$ M, which is based on the hypothesis of the Henry equilibrium of OH radicals between the gas and particle phase (Sander, 2015; Shen et al., 2018). Oxalate can be considered as a tracer for this aqueous chemistry, since it does not have any other significant chemical sources.

Oxalate was detected in the particle-phase samples by ion chromatography. The modeling results of oxalate concentration agreed well with the measured values, and their deviations were in the considered range (Fig. S7). Meanwhile, we can estimate the distribution of major products in irreversible glyoxal/methylglyoxal-OH radical chemistry under different RH conditions, as illustrated in Figure 3d. Generally, oxalate is the major product in wet aerosols, contributing ~60%, and its proportion increases with increasing RH. Besides oxalate, oligomers also play significant roles in glyoxal/methylglyoxal-OH radical chemistry with a contribution of ~30%, and their proportion is maximum under relatively low RH conditions. The RH dependence of the product distribution could mainly be attributed to the particulate concentration of glyoxal/methylglyoxal, which significantly affects the OH radical chemistry. With relatively high carbonyl concentrations (0.1–10 M) in aerosol liquid water, self-reactions of organic molecules become more favorable, resulting in new carbon–carbon bonds and high molecular weight oligomers via radical–radical chemistry (Lim et al., 2013). Moreover, besides OH radical chemistry, reaction with sulfate and ammonium also contribute to the oligomer formation and irreversible uptake of gaseous dicarbonyls (Ortiz-Montalvo et al., 2014; Lim et al., 2016; Lin et al., 2015). The oligomer proportion could be more than 30% in concentrated carbonyl solutions (~0.1 M) and only account for 1% in diluted solutions (~0.01 M).

### 3.4 Relative importance of two partitioning pathways

Table 4 summarizes the particulate concentration of glyoxal and methylglyoxal via reversible and irreversible pathways in
different seasons. The average particulate concentrations of glyoxal (0.43 ng/μg in the reversible pathway and 24.26 ng/μg in the irreversible pathway) were generally higher than those of methylglyoxal (0.25 ng/μg in the reversible pathway and 16.53 ng/μg in the irreversible pathway), mainly due to the relatively higher water solubility and reactivity of glyoxal. Comparing two gas-particle partitioning processes, the irreversible pathway played extremely dominant roles and generally accounted for 96.7% and 95.0% for glyoxal and methylglyoxal, respectively. The proportion of the irreversible pathway varied from 90% to 99.9% and reached its highest in summer for glyoxal (98.8%) and in autumn for methylglyoxal (99.2%), while it was minimum in winter (92.9% for glyoxal and 92.8% for methylglyoxal). Overall, the irreversible pathway played a dominant role in the gas-particle partitioning process for both glyoxal and methylglyoxal in the real atmosphere, while the reversible pathway was also substantial and nonnegligible, especially in winter, with an proportion of ~10%. Furthermore, as discussed above, the particulate concentrations of dicarbonyls and their relative importance were influenced by environmental factors such as relative humidity and particle composition, which could jointly influence both the reversible and irreversible pathways of dicarbonyls. As shown in Figure 4, the proportion of irreversible pathways in the gas-particle partitioning process for dicarbonyls increased with aqueous SNA concentrations, and reached maximum when SNA concentrations were more than 100 M under low RH conditions.

Comprehensively considering the contribution of two pathways in partitioning processes could be conductive to ambient dicarbonyls simulations. Ling et al. (2020) found that the observation and simulation of the gas-phase concentration level of dicarbonyls could reach reasonable agreement when the irreversible uptake and reversible partitioning were incorporated into the model, as these jointly contribute ~62% to the sink of dicarbonyls. Moreover, the contribution of gas-particle partitioning processes of dicarbonyls to SOA formation were higher as the two partitioning pathways were jointly considered. In this study, gas-particle partitioning processes of dicarbonyls accounted for a relatively large proportion of total particle mass (PM$_{2.5}$), on the average of ~5% considering both reversible and irreversible gas-particle partitioning pathways. Since a large fraction of PM$_{2.5}$ mass in Beijing consists of SOAs (~30%) (Huang et al., 2014), we could roughly estimate the contribution of gas-particle partitioning processes of dicarbonyls to SOA yields (by mass). There were approximately 25% SOAs formed from glyoxal and methylglyoxal in this study. However, the particulate dicarbonyls calculated here only contained simple reversible pathways and irreversible pathways driven by OH radicals. More complicated chemical processes like NO$_3$ radical chemistry were not considered, which still resulted in the underestimation of their contribution to SOA formation.

4 Conclusions

We simultaneously measured glyoxal and methylglyoxal concentration in the gas and particle phase in different seasons over urban Beijing. Based on field-measured data, the field-derived gas-particle partitioning coefficients were calculated and found
to be 5–7 magnitudes higher than the theoretical values. Such a large discrepancy provides field evidence that the gas-particle partitioning process does not occur by physical absorption alone but also results from the combined and simultaneous effects of reversible and irreversible pathways. Hydration and oligomerization occurred in the reversible pathway, producing compounds with lower volatility in the condensed phase, and the irreversible pathway could accelerate the uptake of gaseous dicarbonyls. The two pathways jointly contributed to the underestimation of gas-particle partitioning of dicarbonyls. This study systemically considers both reversible and irreversible pathways in the ambient atmosphere for the first time.

Compared to the reversible pathways, the irreversible pathways play a dominant role in the gas-particle partitioning process for dicarbonyls, accounting for ~90% of this process. We recommend the irreversible reactive uptake coefficient for glyoxal and methylglyoxal in different seasons in the real atmosphere. The values we calculated here are higher than those used in model simulations to date, especially for methylglyoxal which exhibits an unexpected salting-in effect under an atmospheric-relevant concentration. We expect the application of these parameterizations will increase the calculated contribution of irreversible uptake of dicarbonyls to SOA formation and narrow the gap between model predictions and field measurements of ambient dicarbonyl concentrations. Moreover, relative humidity and inorganic particle compositions are defined as the most important factor influencing particulate concentration and product distribution of dicarbonyls via both reversible and irreversible pathways, implying the significance of considering different RH conditions in dicarbonyl SOA simulations. Furthermore, we note that there may be other potential explanations for the increase in particulate concentrations and the uncertainty in the gas-particle partitioning process. Physical adsorption of dicarbonyls could be enhanced by water-soluble organics and mineral dust. Other reversible pathways, like adducts formed from glyoxal with inorganic species, like sulfate and ammonia, could also promote the gas-particle partitioning process. Irreversible pathways driven by other oxidants, like NO₃ radicals, can also perform a substantial role. Shen et al. (2016) found that glyoxal could irreversibly produce formic acid, glycolic acid, and oligomers on particles without illumination or extra oxidants. Besides gas-particle partitioning, particulate dicarbonyls formed via the heterogeneous reaction of VOCs could contribute to the uncertainty in partitioning research. Dong et al. (2021) recently revealed that aqueous photooxidation of toluene could yield glyoxal and methylglyoxal via a ring-cleavage process. Overall, the real gas-particle partitioning process of glyoxal and methylglyoxal is more complicated and their contribution to SOA formation is still indistinct; thus, more laboratory experiments and field measurements are urgently needed to improve our understanding of the gas-particle partitioning process for glyoxal and methylglyoxal.

Data availability: The data are accessible by contacting the corresponding author (zmchen@pku.edu.cn).

Author contributions. In the framework of the five field measurements in different seasons, ZC and JH designed the study, and JH performed all carbonyl measurements used in this study, analyzed the data, and wrote the paper. ZC helped interpret the
results, guided the writing, and modified the manuscript. XQ and PD contributed to the methods of sampling and analyzing gas- and particle-phase carbonyls. All authors discussed the results and contributed to the final paper.

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

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Figure 1: Time series of meteorological parameters and gas- and particle-phase glyoxal and methylglyoxal observed in different seasons: (a) summer, 2019.07.20-2019.08.04; (b) winter, 2020.01.05-2020.01.19; (c) autumn, 2020.10.24-2020.11.07; (d) winter, 2021.01.08-2021.01.26; (e) spring, 2021.03.26-2021.04.06. The lines and solid circles indicate the gaseous carbonyls (ppbv) and particulate carbonyls (ng/m³), respectively.
Figure 2: Gas-particle partitioning of dicarbonyls via reversible pathways. (a) The RH dependence of total particulate dicarbonyl concentrations and their pH dependence under high RH conditions. (b) The product distribution for (i) glyoxal and (ii) methylglyoxal under different RH conditions. (c) The gas-particle partitioning coefficients for (i) glyoxal and (ii) methylglyoxal. The black, dark gray, and light gray circles refer to field-measured values, estimated values by the proposed mechanism, and theoretical values calculated by Pankow’s absorptive model, respectively.
Figure 3: Gas-particle partitioning of dicarbonyls via irreversible pathways. (a) The RH dependence of irreversible uptake rate for glyoxal and methylglyoxal. (b) The SNA dependence of uptake coefficients for (i) glyoxal and (ii) methylglyoxal. (c) The correlation between oxalate concentrations measured by ion chromatography (gray lines) and those calculated by modeling coupled with the full kinetic of glyoxal/methylglyoxal + OH (black lines). (d) The corresponding modeled product distribution and O/C (on the right-hand y-axis) in wet aerosols under different RH conditions.
Figure 4: Correlation between the proportion of the irreversible pathway in gas-particle partitioning process for dicarbonyls and aqueous sulfate, nitrate, and ammonia (SNA) concentration in ambient aerosols under different relative humidity conditions.
Table 1: Statistical data for the α-dicarbonyls in gas and particle phase in different seasons.

| Season | Gas phase (ppbv) | Particle phase (ng m$^{-3}$) |
|--------|------------------|-----------------------------|
|        | GL (ppbv) | MG (ppbv) | GL:MG | GL (ng m$^{-3}$) | MG (ng m$^{-3}$) | GL:MG |
| summer | 0.13 ± 0.07 | 0.87 ± 0.54 | 0.15 | 10.18 ± 6.63 | 9.50 ± 5.62 | 1.07 |
| spring | 0.02 ± 0.02 | 0.12 ± 0.08 | 0.17 | 15.24 ± 17.50 | 6.07 ± 2.79 | 2.51 |
| autumn | 0.07 ± 0.03 | 0.15 ± 0.09 | 0.47 | 9.33 ± 4.24 | 9.15 ± 3.62 | 1.02 |
| winter | 0.06 ± 0.05 | 0.11 ± 0.09 | 0.54 | 28.77 ± 25.33 | 14.61 ± 10.15 | 1.97 |
Table 2: Comparison of the field-measured partitioning coefficient $K_f$ values for the dicarbonyls and their corresponding theoretical $K_t$ values in different seasons.

| Coefficients | Dicarbonyl | Season | $K_f$ Average | $K_f$ Range | $K_f$ | $K_f/K_t$ |
|--------------|------------|--------|--------------|-------------|------|---------|
| Glyoxal      |            | summer | $8.11 \times 10^{-4}$ | $(1.25-58.6) \times 10^{-4}$ | $3.27 \times 10^{-10}$ | $2.48 \times 10^{6}$ |
|              |            | autumn | $2.14 \times 10^{-3}$ | $(2.61-224) \times 10^{-4}$ | $6.27 \times 10^{-10}$ | $3.41 \times 10^{6}$ |
|              |            | spring | $1.43 \times 10^{-2}$ | $(0.08-14.6) \times 10^{-2}$ | $5.59 \times 10^{-10}$ | $3.55 \times 10^{7}$ |
|              |            | winter | $1.30 \times 10^{-2}$ | $(0.067-44.2) \times 10^{-2}$ | $1.27 \times 10^{-9}$ | $1.02 \times 10^{7}$ |
| Methyglyoxal |            | summer | $1.49 \times 10^{-4}$ | $(0.833-169) \times 10^{-5}$ | $7.10 \times 10^{-10}$ | $2.10 \times 10^{5}$ |
|              |            | autumn | $9.55 \times 10^{-4}$ | $(0.65-86.9) \times 10^{-4}$ | $1.35 \times 10^{-9}$ | $7.07 \times 10^{5}$ |
|              |            | spring | $1.06 \times 10^{-3}$ | $(0.42-108) \times 10^{-4}$ | $1.21 \times 10^{-9}$ | $8.77 \times 10^{5}$ |
|              |            | winter | $2.60 \times 10^{-3}$ | $(0.34-410) \times 10^{-4}$ | $2.72 \times 10^{-9}$ | $9.93 \times 10^{5}$ |
| Glyoxal      |            | summer | $1.96 \times 10^{8}$ | $(1.71-167) \times 10^{7}$ | $3.29 \times 10^{5}$ | $6.11 \times 10^{2}$ |
|              |            | autumn | $5.08 \times 10^{8}$ | $(1.88-10.2) \times 10^{8}$ | $1.14 \times 10^{6}$ | $3.63 \times 10^{3}$ |
|              |            | spring | $2.53 \times 10^{9}$ | $(1.23-139) \times 10^{8}$ | $9.03 \times 10^{4}$ | $1.92 \times 10^{4}$ |
|              |            | winter | $1.04 \times 10^{9}$ | $(1.37-55.4) \times 10^{8}$ | $4.15 \times 10^{4}$ | $2.55 \times 10^{3}$ |
| Methyglyoxal |            | summer | $4.92 \times 10^{7}$ | $(1.70-363) \times 10^{6}$ | $2.73 \times 10^{3}$ | $1.88 \times 10^{4}$ |
|              |            | autumn | $8.52 \times 10^{7}$ | $(3.66-15.7) \times 10^{7}$ | $9.50 \times 10^{3}$ | $2.00 \times 10^{5}$ |
|              |            | spring | $1.33 \times 10^{8}$ | $(5.22-456) \times 10^{6}$ | $7.49 \times 10^{3}$ | $1.36 \times 10^{5}$ |
|              |            | winter | $2.63 \times 10^{8}$ | $(1.03-175) \times 10^{7}$ | $3.44 \times 10^{4}$ | $9.01 \times 10^{4}$ |
Table 3: Summary of calculated uptake coefficients $\gamma$ and effective uptake rate coefficient $k_{\text{eff, uptake}}$ in different seasons for glyoxal and methylglyoxal.

| Dicarbonyl | Season | T (K) | RH (%) | $\gamma$ ($\times 10^{-3}$) | $k_{\text{eff, uptake}}$ (s$^{-1}$) |
|------------|--------|-------|--------|-----------------------------|----------------------------------|
|            |        |       | Average | Min | Max | ($\times 10^{-4}$) |
| GL         | Summer | 301.1 | 67.7   | 4.15 | 0.12 | 7.30 | 1.61 |
|            | Autumn | 287.2 | 45.4   | 8.62 | 0.29 | 12.9 | 4.83 |
|            | Spring | 289.4 | 54.0   | 11.7 | 1.24 | 14.9 | 6.85 |
|            | Winter | 273.5 | 54.0   | 10.6 | 0.56 | 14.4 | 3.59 |
| *General   |        | 287.8 | 59.0   | 8.0  | 0.46 | 11.4 | 3.38 |
| MG         | Summer | 301.1 | 67.7   | 1.01 | 0.02 | 2.09 | 0.25 |
|            | Autumn | 287.2 | 45.4   | 1.83 | 0.04 | 3.94 | 0.92 |
|            | Spring | 289.4 | 54.0   | 2.36 | 0.06 | 4.34 | 0.69 |
|            | Winter | 273.5 | 54.0   | 3.45 | 0.11 | 5.83 | 0.77 |
| *General   |        | 287.8 | 59.0   | 2.0  | 0.05 | 3.8  | 0.55 |

*General is the average value of all the samples observed in the five field observations.
Table 4: Calculated relative importance of reversible and irreversible pathways in the gas-particle partitioning processes and their contribution to the particle matter.

| Season | Glyoxal         | Methylglyoxal | Contribution to particulate matters |
|--------|-----------------|---------------|-------------------------------------|
|        | \(a [X]_{\text{rev}}\) | \(b [X]_{\text{irrev}}\) | \(a [X]_{\text{rev}}\) | \(b [X]_{\text{irrev}}\) |                          |
| Summer | 0.17 (1.2%)     | 18.87 (98.8%) | 0.25 (5.5%)                       | 20.55 (94.5%) | 3.98%                   |
| Autumn | 0.14 (0.7%)     | 23.91 (99.3%)| 0.12 (0.8%)                       | 17.02 (99.2%)| 4.12%                   |
| Spring | 0.26 (1.9%)     | 15.94 (98.1%)| 0.09 (1.5%)                       | 14.16 (98.5%)| 3.05%                   |
| Winter | 0.89 (7.1%)     | 34.70 (92.9%)| 0.38 (7.2%)                       | 12.59 (92.8%)| 4.86%                   |
| General| 0.43 (3.3%)     | 24.26 (96.7%)| 0.25 (5.0%)                       | 16.53 (95.0%)| 4.15%                   |

\(a [X]_{\text{rev}}\) is the concentration of particle-phase carbonyl via reversible pathway (ng·μg\(^{-1}\)) and its proportion (%).

\(b [X]_{\text{irrev}}\) is the concentration of particle-phase carbonyl via irreversible pathway (ng·μg\(^{-1}\)) and its proportion (%).

General is the average value of all the samples observed in the five field observations.