1	Water adsorption and hygroscopic growth of six anemophilous pollen species: the
2	effect of temperature
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## 24 Abstract

Hygroscopicity largely affects environmental and climatic impacts of pollen grains, one 25 important type of primary biological aerosol particles in the troposphere. However, our knowledge 26 in pollen hygroscopicity is rather limited, and especially the effect of temperature has rarely been 27 explored before. In this work three different techniques, including a vapor sorption analyzer, 28 diffusion reflectance infrared Fourier transform spectroscopy (DRIFTS) and transmission Fourier 29 transform infrared spectroscopy (transmission FTIR) were employed to characterize six 30 anemophilous pollen species and to investigate their hygroscopic properties as a function of 31 relative humidity (RH, up to 95%) and temperature (5 or 15, 25 and 37 °C). Substantial mass 32 increase due to water uptake was observed for all the six pollen species, and at 25 °C the relative 33 mass increase at 90% RH, when compared to that at <1% RH, ranged from  $\sim30$  to  $\sim50\%$ , varying 34 with pollen species. It was found that the modified  $\kappa$ -Köhler equation can well approximate mass 35 hygroscopic growth of all the six pollen species, and the single hygroscopicity parameter ( $\kappa$ ) was 36 determined to be in the range of 0.034±0.001 to 0.061±0.007 at 25 °C. In-situ DRIFTS 37 measurements suggested that water adsorption by pollen species was mainly contributed by OH 38 groups of organic compounds they contained, and good correlations were indeed found between 39 hygroscopicity of pollen species and the amount of OH groups, as determined using transmission 40 FTIR. Increase in temperature would in general lead to decrease in hygroscopicity, except for 41 pecan pollen. For example,  $\kappa$  values decreased from 0.073±0.006 at 5 °C to 0.061±0.007 at 25 °C 42 and to  $0.057\pm0.004$  at 37 °C for populus tremuloides pollen, and decreased from  $0.060\pm0.001$  at 43 15 °C to 0.054±0.001 at 25 °C to 0.050±0.002 at 37 °C for paper mulberry pollen. 44

## 46 **1 Introduction**

Primary biological aerosol particles (PBAPs), an important type of aerosol particles in the 47 48 troposphere, are directly emitted from the biosphere and include pollen, fungal spores, bacteria, viruses, algae, and so on (Després et al., 2012; Fröhlich-Nowoisky et al., 2016). Emission and 49 abundance of PBAPs are quite uncertain, and annual emission fluxes are estimated to be in the 50 range of <10 to ~1000 Tg for total PBAPs and 47-84 Tg for pollen (Després et al., 2012). Pollen, 51 and PBAPs in general, are of great concerns due to their various impacts on the Earth system (Sun 52 and Ariya, 2006; Ariya et al., 2009; Georgakopoulos et al., 2009; Morris et al., 2011; Morris et al., 53 54 2014; Fröhlich-Nowoisky et al., 2016). For example, they can be allergenic, infectious or even toxic, affecting the health of human and other species in the ecological systems over different 55 scales (Douwes et al., 2003; Reinmuth-Selzle et al., 2017; Shiraiwa et al., 2017). The geographical 56 dispersion of anemophilous plants largely relies on pollen dispersal, which in turn depends on the 57 emission, transport and deposition of pollen grains; therefore, pollen plays a key role in the 58 59 evolution of many ecosystems (Womack et al., 2010; Fröhlich-Nowoisky et al., 2016). In addition, PBAPs can serve as giant cloud condensation nuclei (CCN) and ice nucleating particles (INPs), 60 significantly impacting the formation and properties of clouds and thus radiative balance and 61 precipitation (Möhler et al., 2007; Ariya et al., 2009; Pratt et al., 2009; Pope, 2010; Pummer et al., 62 2012; Gute and Abbatt, 2018). It has also been proposed that PBAPs may have significant impacts 63 on chemical composition of aerosol particles via heterogeneous and multiphase chemistry 64 (Deguillaume et al., 2008; Estillore et al., 2016; Reinmuth-Selzle et al., 2017; Shiraiwa et al., 2017). 65 Hygroscopicity is one of the most important physicochemical properties of pollen (as well 66 as aerosol particles in general). Hygroscopicity largely impacts the transport and deposition of 67 pollen grains (Sofiev et al., 2006), therefore affecting their lifetimes, abundance and 68

spatiotemporal distribution. In addition, hygroscopicity is closely linked to the ability of aerosol 69 particles to serve as CCN and INPs (Petters and Kreidenweis, 2007; Kreidenweis and Asa-Awuku, 70 2014; Laaksonen et al., 2016; Tang et al., 2016). Several previous studies have measured the 71 hygroscopicity and CCN activities of pollen (Diehl et al., 2001; Pope, 2010; Griffiths et al., 2012; 72 Lin et al., 2015; Steiner et al., 2015; Prisle et al., 2018) and other PBAPs such as bacteria (Pasanen 73 74 et al., 1991; Reponen et al., 1996; Franc and DeMott, 1998; Ko et al., 2000; Lee et al., 2002; Bauer et al., 2003). For example, water uptake of eleven pollen species was studied using an analytical 75 balance (Diehl et al., 2001), and the mass of pollen was found to be increased by 3-16% at 73% 76 77 RH and by  $\sim 100-300\%$  at 95% RH, compared to that at <1% RH. An electrodynamic balance was employed to investigate hygroscopic growth of eight types of pollen (Pope, 2010; Griffiths et al., 78 2012), and it was found that their hygroscopic growth can be approximated by the modified  $\kappa$ -79 Köhler equation, with single hygroscopicity parameters being around 0.1 (depending on the 80 assumed pollen density, as discussed in Section 3.2). 81

82 Previous measurements were mostly carried out at or close to room temperature, and the effects of temperature on hygroscopic properties of pollen and other types of PBAPs are yet to be 83 elucidated. To our knowledge, only one previous study (Bunderson and Levetin, 2015) explored 84 the effect of temperature (4, 15 and 20 °C) on water uptake by Juniperus ashei, Juniperus 85 monosperma and Juniperus pinchotii pollen. It is important to account for the temperature effects, 86 because ambient temperatures range from below -70 to >30 °C in the troposphere. In particular, 87 88 the altitude of 0.5-2.0 km to which pollen can be easily transported (Noh et al., 2013) may have temperatures close to or lower than the chilling temperatures for vegetative species (up to 16.5 °C) 89 90 (Melke, 2015). Moreover, the temperature in the respiratory tract can reach up to 37 °C (the 91 physiological temperature). In the work presented here, a vapor sorption analyzer (VSA) was

employed to investigate hygroscopic growth of pollen grains at different temperature (5 or 15, 25, and 37 °C), a range covering the chilling temperature to the physiological temperature. Water uptake by pollen was also examined using diffusion reflectance infrared Fourier transform spectroscopy at room temperature to complement the VSA results. Furthermore, transmission Fourier transformation infrared spectroscopy was used to characterize functional groups of dry pollen grains, in an attempt to seek potential links between chemical composition of pollen grains and their hygroscopic properties.

## 99 **2 Experimental sections**

Six pollen species, all from anemophilous plants, were investigated in this work, including populus tremuloides and populus deltoides (provided by Sigma Aldrich) as well as ragweed, corn, pecan and paper mulberry (provided by Polysciences, Inc.). The six pollen species were chosen in our work primarily because they were commercially available. Furthermore, these plants are also widely distributed in the globe. For example, corn is the most produced grain in the world (International-Grains-Council, 2019), and up to 50% of pollen-related allergic rhinitis cases in North America are caused by ragweed pollen (Taramarcaz et al., 2005).

## 107 **2.1 Fourier transformation infrared spectroscopy**

The adsorption of water by pollen species were studied using in-situ diffusion reflectance infrared Fourier transform spectroscopy (DRIFTS) at room temperature (~25 °C). This technique was described in details in our previous work (Ma et al., 2010), and similar setups have also been used by other groups to investigate adsorption of water by mineral dust (Joshi et al., 2017; Ibrahim et al., 2018). Infrared spectra were recorded using a Nicolet 6700 Fourier transformation infrared spectrometer (FTIR, Thermo Nicolet Instrument Corporation), equipped with an in-situ diffuse reflection chamber and a high-sensitivity mercury cadmium telluride (MCT) detector cooled by

liquid nitrogen. A pollen sample (about 10 mg for each sample) under investigation was placed 115 into a ceramic crucible which was located in the in-situ chamber. A dry air flow and a humidified 116 air flow were first mixed and then delivered into the chamber, and the total flow rate was set to 117 200 mL/min (standard condition). Relative humidity (RH) in the chamber could be adjusted by 118 varying the flow rate ratio of the dry flow to the humidified flow, and was monitored online using 119 120 a moisture meter (CENTER 314) with an absolute uncertainty of  $\pm 2\%$ . Prior to each experiment, the sample was flushed with dry air for 3 h at 25 °C, and the reference spectrum was recorded after 121 the pretreatment. Infrared spectra were collected and analyzed using OMNIC 6.0 software (Nicolet 122 Corp.). All the spectra reported here were recorded with a wavenumber resolution of 4 cm<sup>-1</sup>, and 123 100 scans were averaged to produce a spectrum. Water adsorption was equilibrated for at least 30 124 min at each RH to ensure that the equilibrium between water vapor and adsorbed water was 125 reached. 126

Pollen samples used in this work were also characterized using transmission FTIR equipped with a deuterated triglycine sulfate detector (DTGS) detector. Pollen grains and KBr were mixed with a mass ratio of approximately 1:100 and ground in an agate mortar, and the mixture was then pressed into a clear disc. Transmission FTIR was employed to examine these discs, and a pure KBr disc was used as the reference. All the spectra, each of which was the average of 100 scans, were also recorded with a wavenumber resolution of 4 cm<sup>-1</sup>.

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### 2.2 Vapor sorption analyzer

Hygroscopic growth of pollen grains was further investigated using a vapor sorption analyzer (Q5000 SA, TA Instruments, New Castle, DE, USA) described in our previous work (Gu et al., 2017; Guo et al., 2018; Jia et al., 2018). In brief, this instrument measured the sample mass as a function of RH under isothermal conditions. The instrument can be operated in the temperature

138 range of 5-85 °C with a temperature accuracy of ±0.1 °C and in the RH range of 0-98 % with an 139 absolute accuracy of  $\pm 1\%$ . RH in the humidity chamber was regulated by using two mass flow controllers to control the dry and humidified nitrogen flows very precisely. The accuracy in RH 140 141 control was routinely checked by measuring the DRH values for a series of standard compounds, e.g., NaCl, (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>, KCl, and etc., and the difference between the measured and theoretical 142 DRH was always <1%. The mass measurement had a range of 0-100 mg and a sensitivity of  $\pm 0.01$ 143  $\mu$ g. The initial mass of each sample used in this work was in the range of 0.5-1 mg. For each of 144 the first three types of pollen species (populus tremuloides, populus deltoides and ragweed pollen), 145 three samples in total were investigated, and each sample was studied under isothermal conditions 146 at 5, 25 and 37 °C. For each of the other three types of pollen species (corn, pecan and paper 147 mulberry pollen), experiments were carried out at 15 °C instead of 5 °C, because the instrument 148 could only be cooled down to 15 °C due to a technical problem after we finished experiments for 149 the first three pollen species. 150



Figure 1. Change of RH (black curve, left *y*-axis) and normalized sample mass (blue curve, right *y*-axis) with time for a typical experiment in which hygroscopic growth of pollen grains was measured. In this figure a dataset for paper mulberry pollen at 25 °C is plotted as an example.

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For the first sample, at each temperature the sample was first dried at 0% RH (the actual 156 RH was measured to be <1%); after that, RH was increased stepwise to 95% with an increment of 157 5% per step and then switched back to <1% to dry the sample again. At each RH, the sample was 158 equilibrated with the environment (i.e. until the sample mass became stable) before RH was 159 160 changed to the next value, and the sample mass was considered to be stabilized when the mass change was <0.05% within 30 min. Such a measurement at one temperature could take several 161 days. In order to reduce experimental time, the second and third samples were investigated in a 162 similar way as the first sample, except that RH was increased stepwise to 90% with an increment 163 of 10% per step. A typical experimental dataset is displayed in Figure 1 as an example to illustrate 164 the change of RH and normalized sample mass with experimental time. 165

# 166 **3 Results and discussion**

### 167 **3.1 FTIR characterization of pollen samples**

### 168 **3.1.1 Infrared spectra of dry pollen samples**

Figure 2 shows the transmission FTIR spectra of the six pollen species investigated in our work, and peak assignments can be found in Table 1. A broad band in the range of 3600-3000 cm<sup>-1</sup>, attributed to O-H stretching vibration (Stuart, 2004; Pummer et al., 2013), and two sharp peaks at 2920 and 2850 cm<sup>-1</sup>, attributed to C-H stretching (Eliason et al., 2003; Stuart, 2004; Pummer et al., 2013), were observed for all the pollen species. The two peaks at 1747 and 1658 cm<sup>-1</sup> were assigned to alkyl ester carbonyls (Pappas et al., 2003; Najera et al., 2009; Pummer et al., 2013), and the two peaks at 1549 and 1458 cm<sup>-1</sup> (1411 cm<sup>-1</sup> for paper mulberry pollen) were assigned to C=C stretching and H-C-H deformation (Stuart, 2004; Pummer et al., 2013). In addition, the three peaks at 1053, 997 and 845 cm<sup>-1</sup> were assigned to C-O stretching, C-C stretching, and C-H out-ofplane bending, respectively (Stuart, 2004; Najera et al., 2009; Pummer et al., 2013).



179

180 **Figure 2.** Transmission FTIR spectra of six pollen species investigated in this work.

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OH groups and C-H groups in organic compounds are generally considered to be 182 hydrophilic and hydrophobic, and one may expect that the amount of OH groups (relative to that 183 of C-H groups) that organic samples contain may affect their hygroscopicity. For example, it was 184 found in many previous studies (Eliason et al., 2003; Asad et al., 2004; Hung et al., 2005; Najera 185 et al., 2009) that heterogeneous reactions of organic materials with O<sub>3</sub> and OH radicals would 186 increase the IR absorption intensity for the O-H stretching mode and decrease the IR absorption 187 intensity for the C-H stretching mode, meanwhile leading to the enhancement in their 188 hygroscopicity. Therefore, in this work we use the intensity ratio of the O-H stretching vibration 189

band (3000-3600 cm<sup>-1</sup>) to the C-H stretching mode (2920 cm<sup>-1</sup>) to qualitatively represent the 190 191 amount of OH groups pollen samples contain. As shown in Figure 2, the six pollen species examined in our work can be roughly classified into two catalogues: 1) for populus deltoides, 192 populus tremuloides and paper mulberry pollen, the O-H stretching vibration band is more 193 intensive than the C-H stretching mode, indicating that they contain relatively high levels of OH 194 195 groups; 2) for ragweed, pecan and corn pollen, the O-H stretching vibration band is less intensive than the C-H stretching mode, indicating that they contain relatively low levels of OH groups. The 196 relation between the amount of OH groups that pollen species contain and their hygroscopicity 197 198 will be further discussed in Section 3.3.

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**Table 1.** Vibrational mode assignment for six pollen species investigated in this work.

wavenumber (cm <sup>-1</sup> )	vibrational mode
3600-3000	O-H stretching
2920 and 2820	C-H stretching
1747 and 1658	alkyl ester carbonyls
1549	C=C stretching
1458 and 1411	H-C-H deformation
1053	C-O stretching
997	C-C stretching
845	C-H out-of-plane bending

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### **3.1.2 Infrared spectra of pollen samples at different RH**

In-situ DRIFTS was employed to explore adsorption of water by pollen grains. Typical spectra of populous deltoides pollen as a function of RH up to 87%, relative to that at <1% RH, are displayed in Figure 3. DRIFTS spectra of other pollen samples at different RH can be found in Figures S1-S5 in the supplement, and are very similar to those for populous deltoides pollen. As

evident from Figure 3, several IR peaks (e.g., 3593, 3205, 2135, and 1616 cm<sup>-1</sup>) appeared in the 207 spectra at elevated RH, when compared with that at <1% RH, and their intensities increased with 208 increasing RH. The peaks at 3205, 2135 and 1616 cm<sup>-1</sup> can be assigned to the stretching, 209 association and bending modes of adsorbed water (Goodman et al., 2001; Schuttlefield et al., 210 2007a; Ma et al., 2010; Hatch et al., 2011; Song and Boily, 2013; Yeşilbaş and Boily, 2016; Joshi 211 et al., 2017; Ibrahim et al., 2018). 212



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Figure 3. In-situ DRIFTS spectra of populous deltoides pollen as a function of RH (<1, 26, 41, 57, 214 71, 81 and 87%) at 25 °C. 215

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222 interacted with OH groups in pollen samples (Iwamoto et al., 2003). These results imply that water adsorption by pollen samples could be mainly contributed by OH groups of organic compounds 223 they contained. Both C-OH and C(O)-OH groups can contribute to water adsorption by pollen 224 samples, though their relative contribution cannot be resolved in our work. In addition, other 225 factors, such as porosity and internal structure, may also be important for hygroscopic properties 226 of pollen grains. The intensities of IR peaks at ~3600 cm<sup>-1</sup> were used to represent the amount of 227 water adsorbed by pollen samples. Table 2 summarizes integrated areas of IR peaks at 3600 cm<sup>-1</sup> 228 as a function of RH for the six pollen species examined in our work, suggesting that the amount 229 230 of adsorbed water by pollen samples increased with RH.

Table 2. Integrated areas of IR peaks (at  $\sim$ 3600 cm<sup>-1</sup>) of adsorbed water as a function of RH for the six pollen species investigated in this work. Wavenumber ranges used for integration are 3750-3300 cm<sup>-1</sup> for populus deltoides pollen, 3750-3350 cm<sup>-1</sup> for populus tremuloides pollen, 3750-3400 cm<sup>-1</sup> for ragweed pollen, 3750-3500 cm<sup>-1</sup> for corn pollen, 3750-3450 cm<sup>-1</sup> for pecan pollen, and 3750-3300 cm<sup>-1</sup> for paper mulberry pollen.

RH (%)	peak area	RH (%)	peak area	RH (%)	peak area
populus deltoides		populus tremuloides		ragweed	
<1	0	<1	0	<1	0
26	22.7	24	5.5	26	10.1
41	36.9	41	16.4	42	18.9
57	57.4	56	35.4	50	24.5
71	93.6	70	66.5	56	30.2
79	137.6	78	91.2	69	49.7
81	164.7	87	156.9	88	104.6
87	293.1				
corn		pecan		paper mulberry	
<1	0	<1	0	<1	0

26	10.0	26	8.6	26	10.2
42	21.5	43	16.9	43	17.7
58	41.9	58	29.5	51	23.1
73	87.5	73	60.0	59	29.8
89	222.2	89	338.9	71	46.7
				86	105.1

# 238 **3.2 Mass hygroscopic growth**

239 **3.2.1 Hygroscopicity parameterizations** 

The single hygroscopicity parameter,  $\kappa$ , is widely used to describe the hygroscopicity of aerosol particles under both subsaturation and supersaturation (Petters and Kreidenweis, 2007). When the Kelvin effect is negligible (this is valid for pollen grains which are typically >1 µm), the dependence of diameter-based growth factor (GF) on RH can be linked to  $\kappa$  via Eq. (1) (Petters and Kreidenweis, 2007; Tang et al., 2016):

$$RH = \frac{GF^3 - 1}{GF^3 - 1 + \kappa} \quad (1)$$

## If we further assume that the particle is spherical, Eq. (1) can be transformed to Eq. (2):

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$$\frac{1}{RH} = 1 + \frac{\kappa}{GF^3 - 1} = 1 + \frac{\kappa}{\frac{V}{V_0} - 1} = 1 + \kappa \frac{V_0}{V - V_0} = 1 + \kappa \frac{V_0}{V_w}$$
(2)

where V,  $V_0$  and  $V_w$  are the volumes of the particle at the given RH, the dry particle and water associated with the particle at the given RH. In order for Eq. (2) to be valid, it is also assumed that at a given RH, V is equal to the sum of  $V_0$  and  $V_w$ . Eq. (2) can be further transformed to Eqs. (3-4):

$$\frac{1}{RH} = 1 + \kappa \frac{\rho_w}{\rho_p} \frac{m_0}{m_w} \quad (3)$$

252 
$$\frac{m_w}{m_0} = \kappa \cdot \frac{\rho_w}{\rho_p} / (\frac{1}{RH} - 1) \quad (4)$$

where  $\rho_{\rm w}$  and  $\rho_{\rm p}$  are the density of water and the dry particle, and  $m_0$  and  $m_{\rm w}$  are the mass of the dry particle and water associated with the particle at the given RH. Since the particle mass, *m*, is equal to the sum of  $m_0$  and  $m_{\rm w}$ , Eq. (5) can be derived from Eq. (4):

256 
$$\frac{m}{m_0} = 1 + \kappa \frac{\rho_W}{\rho_p} / (\frac{1}{RH} - 1) \quad (5)$$

257 Using an electrodynamic balance, Pope and co-workers (Pope, 2010; Griffiths et al., 2012) measured hygroscopic growth of eight types of pollen grains, and found that their mass change 258 with RH can be approximated by Eq. (5). It should be noted that the original equation derived by 259 Pope and co-workers (Pope, 2010; Griffiths et al., 2012) has a different format from but is 260 essentially equivalent to Eq. (5). Eq. (5) relates mass growth experimentally measured in our work 261 to the single hygroscopicity parameter ( $\kappa$ ), which has been widely used in atmospheric science to 262 describe hygroscopic properties of aerosol particles under subsaturation as well as their CCN 263 activities under supersaturation; nevertheless, a few assumptions are needed to derive Eq. (5), as 264 265 discussed.

The Freundlich adsorption isotherm is another widely used equation to describe the change of sample mass with RH due to water uptake (Atkins, 1998; Skopp, 2009; Hatch et al., 2011; Tang et al., 2016):

$$\frac{m}{m_0} = 1 + A_f \cdot \sqrt[B_f]{RH} \quad (6)$$

where  $A_f$  and  $B_f$  are empirical Freundlich constants related to the adsorption capacity and strength. One advantage of the Freundlich adsorption isotherm is that it provides a direct relationship between RH and mass growth which was experimentally measured in our work, without any additional assumptions. In addition, the BET (Brunauer-Emmett-Teller) adsorption isotherm is also widely used to describe water adsorption by insoluble solid particles (Brunauer et al., 1938; Goodman et al., 2001; Henson, 2007; Ma et al., 2010; Tang et al., 2016; Joshi et al., 2017). While the BET adsorption isotherm typically works well for water adsorption of a few monolayers, the mass of adsorbed water, as shown in Section 3.2.2, can reach up to 50% of the dry pollen mass at high RH; therefore, in this work we did not attempt to use the BET adsorption isotherm to describe water adsorption by pollen grains. Another reason that we did not attempt to use the BET adsorption isotherm is that the BET adsorption isotherm is mathematically more complex and requires the BET surface area to be known.

## 282 **3.2.2 Mass hygroscopic growth at room temperature**

Figure 4 displays the sample mass (normalized to that at 0% RH) as a function of RH for pecan pollen at 25 °C. Significant increase in sample mass was observed at elevated RH due to uptake of water. Compared to that at <1% RH, the sample mass increased by  $(2.3\pm0.3)$ % at 30% RH,  $(6.4\pm0.2)$ % at 60% RH,  $(30.3\pm0.4)$ % at 90% RH, and up to ~72% at 95% RH. As shown by the data compiled in Tables S1-S3 in the supplement, substantial increases in sample mass were also observed for the other five types of pollen species at 25 °C (as well as 5 and 37 °C).



Figure 4. Measured change of sample mass (normalized to that at dry conditions, i.e.  $m/m_0$ ) of pecan pollen as a function of RH (0-95%) at 25 °C. The experimental data are fitted with the modified  $\kappa$ -Köhler equation (solid red curve) and the Freundlich adsorption isotherm (dashed blue curve).

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Hygroscopic properties exhibited considerable variations among different pollen species. 295 Figure 5a compares the measured ratios of sample mass at 90% RH to that at <1% RH,  $m(90\%)/m_0$ , 296 for the six pollen species investigated in this work. We specifically discuss mass changes of pollen 297 grains at 90% RH (relative to that at <1% RH) because aerosol hygroscopic growth at 90% RH 298 was widely reported by laboratory and field studies (Kreidenweis and Asa-Awuku, 2014). As 299 shown in Figure 5a,  $m(90\%)/m_0$  determined at 25 °C ranged from 1.293±0.028 (ragweed pollen) 300 301 to 1.476±0.094 (populus deltoides pollen), i.e. the amount of water adsorbed/absorbed by the six different pollen species at 90% RH varied between ~30% to ~50% of the dry mass. 302



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Figure 5. Measured ratios of sample mass at 90% RH to that at <1% RH (a) and derived  $\kappa$  values (b) for six pollen species at different temperatures. The lowest temperatures were 5 °C for populus deltoides (populus del.), populus tremuloides (populus tre.) and ragweed pollen, and 15 °C for corn, pecan and paper mulberry pollen.

As shown in Figure 4, the increase of pecan pollen mass with RH at 25 °C could be satisfactorily described by the modified  $\kappa$ -Köhler equation for the entire RH range (up to 95%). On the contrary, the Freundlich adsorption isotherm significantly underestimated the sample mass at low RH, although it represented the experimental data at high RH reasonably well. In addition, we found that the modified  $\kappa$ -Köhler equation could also approximate the dependence of sample mass on RH for all the six types of pollen species investigated in this work at different temperatures. If we use Eq. (5) to fit  $m/m_0$  against RH,  $\kappa \cdot \rho_w / \rho_p$  can be derived. The bulk densities of dry pollen grains were found to vary with species but typically fall into the range of 0.5-2 g cm<sup>-3</sup> (Harrington and Metzger, 1963; Hirose and Osada, 2016), and for simplicity  $\rho_p$  was assumed to be 1 g cm<sup>-3</sup> in this work (i.e.  $\rho_w / \rho_p$  is equal to 1). With the assumptions on dry particle density and also particle sphericity,  $\kappa$  could then be derived from the measured RH-dependent sample mass at a given temperature.

Table 3 summarizes the average  $\kappa$  values at different temperatures for the six pollen species 321 investigated in this work. At 25 °C, the  $\kappa$  values were found to increase from 0.034±0.001 for 322 323 ragweed pollen to 0.061±0.007 for populus tremuloides pollen, varied by almost a factor of 2. The  $\kappa$  values measured by Pope and co-workers (Pope, 2010; Griffiths et al., 2012) were approximately 324 in the range of 0.05-0.11 (assuming that  $\rho_w/\rho_p$  is equal to 1), in reasonably good agreement with 325 these reported in our work. It should be noted that in order to convert the measured mass growth 326 to diameter growth and  $\kappa$  values, one key assumption is particle sphericity; nevertheless, pollen 327 grains are known to be non-spherical and porous, and therefore our derived  $\kappa$  values might be 328 smaller than the actual values. For example, although the mass increase was substantial (around 329 30-50 % at 90% RH) for the six pollen species examined, their  $\kappa$  values at 25 °C were derived to 330 331 be in the range of 0.034-0.061, significantly smaller than those (0.1-0.2) for typical secondary organic aerosols produced in smog chamber studies (Petters and Kreidenweis, 2007; Kreidenweis 332 333 and Asa-Awuku, 2014).

Table 3. Single hygroscopicity parameters ( $\kappa$ ) derived in this work for the six pollen species at different temperatures. All the errors given in this work are standard deviations.

pollen type	<i>T</i> (°C)	sample 1	sample 2	sample 3	average
	5	$0.071 \pm 0.001$	$0.078 \pm 0.001$	$0.057 \pm 0.002$	$0.069 \pm 0.011$

populus	25	$0.054 \pm 0.001$	$0.067 \pm 0.002$	$0.049 \pm 0.002$	$0.057 \pm 0.009$
deltoides	37	$0.058 \pm 0.002$	$0.051 \pm 0.001$	$0.045 \pm 0.002$	$0.051 \pm 0.007$
populus	5	0.068±0.001	0.073±0.001	$0.079 \pm 0.001$	0.073±0.006
tremuloides	25	$0.053 \pm 0.002$	$0.063 \pm 0.002$	$0.066 \pm 0.002$	$0.061 \pm 0.007$
	37	$0.052 \pm 0.002$	$0.059 \pm 0.002$	$0.060 \pm 0.002$	$0.057 \pm 0.004$
ragweed	5	0.042±0.001	0.037±0.002		$0.040 \pm 0.004$
	25	$0.033 \pm 0.002$	$0.034 \pm 0.003$	$0.034 \pm 0.002$	$0.034 \pm 0.001$
	37	$0.027 \pm 0.001$	$0.031 \pm 0.002$	$0.031 \pm 0.002$	$0.030 \pm 0.002$
corn	15	0.051±0.001	$0.059 \pm 0.002$	$0.055 \pm 0.002$	$0.055 \pm 0.004$
	25	$0.046 \pm 0.002$	$0.053 \pm 0.002$	$0.054 \pm 0.002$	$0.051 \pm 0.004$
	37	$0.041 \pm 0.002$	$0.048 \pm 0.002$	$0.048 \pm 0.002$	$0.046 \pm 0.004$
pecan	15	0.049±0.001	0.038±0.001	$0.039 \pm 0.001$	$0.042 \pm 0.006$
	25	$0.036 \pm 0.001$	$0.034 \pm 0.001$	$0.035 \pm 0.001$	$0.035 \pm 0.001$
	37	$0.033 \pm 0.001$	$0.035 \pm 0.002$	$0.037 \pm 0.001$	$0.035 \pm 0.002$
paper	15	$0.059 \pm 0.002$	$0.060 \pm 0.002$	$0.060 \pm 0.002$	$0.060 \pm 0.001$
mulberry	25	$0.054 \pm 0.001$	$0.054 \pm 0.001$	$0.054 \pm 0.001$	$0.054 \pm 0.001$
	37	$0.048 \pm 0.002$	$0.050 \pm 0.002$	$0.051 \pm 0.002$	$0.050 \pm 0.002$

### 338 **3.3 Discussion**

### 339 **3.3.1 Reconciliation between IR and VSA results**

Our in-situ DRIFTS measurements, as discussed in Section 3.1.2, suggested that water 340 uptake by pollen samples was mainly contributed by OH groups of organic compounds they 341 contained; therefore, one may expect that pollen species which contain higher levels of OH groups 342 would exhibit higher hygroscopicity. Transmission FTIR characterization of pollen species 343 344 (Section 3.1.1) showed that populus deltoides, populus tremuloides and paper mulberry pollen contained relatively high levels of OH groups, and indeed their hygroscopicity ( $\kappa$ : 0.053-0.054 at 345 25 °C) was higher than the other three pollen species, as shown in Figure 5 and Table 3. For 346 347 comparison, ragweed and pecan pollen contained relatively low levels of OH groups and correspondingly exhibited lower hygroscopicity ( $\kappa$ : 0.033-0.036 at 25 °C). Corn pollen appeared 348

to be an exception: it contained relatively low levels of OH groups but displayed medium hygroscopicity ( $\kappa$ : ~0.046 at 25 °C). As a result, our results may imply that in addition to chemical composition, other physicochemical properties, such as porosity and internal structure of pollen grains, could also play an important role in determining the hygroscopicity of pollen species. One clue came from environmental scanning electron microscopy observations (Pope, 2010), revealing that pollen grains started to swell internally before significant water uptake on the surface took place.

In our work two complementary techniques were employed to explore hygroscopic 356 properties of pollen species. VSA measured the amount of water absorbed/adsorbed by pollen 357 grains as a function of RH in a quantitative manner, whereas the intensities of IR peaks of adsorbed 358 water at different RH, as characterized by DRIFTS, can be used semi-quantitatively to represent 359 the amount of water associated with particles (Goodman et al., 2001; Schuttlefield et al., 2007b; 360 Ma et al., 2010; Yeşilbaş and Boily, 2016; Joshi et al., 2017; Ibrahim et al., 2018). We compare 361 our VSA results (i.e. the relative mass change due to water uptake) to the DRIFTS results (i.e. 362 integrated area of IR peaks at ~3600 cm<sup>-1</sup>). As shown in Figure 6, good correlations between VSA 363 and DRIFTS results are found for all the six pollen species, suggesting that DRIFTS can be used 364 365 to represent the amount of adsorbed water, at least in a semi-quantitative manner.



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Figure 6. Integrated areas of IR peaks at ~3600 cm<sup>-1</sup> versus relative mass increase due to water uptake,  $(m-m_0)/m_0$ , for six pollen species: (a) populus deltoides; (b) populus tremuloides; (c) ragweed; (d) corn; (e) pecan; (f) paper mulberry.

## 371 **3.3.2 Effect of temperature**

Figure 5a shows the comparison of the measured ratios of sample mass at 90% RH to that at <1% RH,  $m(90\%)/m_0$ , at different temperatures for the six pollen species. It can be concluded from Figure 5a that except for pecan pollen for which a small increase in  $m(90\%)/m_0$  occurred when temperature increased from 25 to 37 °C, increase in temperature would lead to small but nevertheless significant decrease in  $m(90\%)/m_0$ . For example,  $m(90\%)/m_0$  decreased from 1.597±0.100 at 5 °C to 1.476±0.094 at 25 °C and to 1.427±0.060 at 37 °C for populus deltoides pollen, and from 1.338±0.036 at 5 °C to 1.293±0.028 at 25 °C and to 1.249±0.002 at 37 °C for
ragweed pollen.

We further derived  $\kappa$  values at different temperatures for the six pollen species, and the results are plotted in Figure 5b and summarized in Table 3. Increase in temperature would lead to decrease in  $\kappa$  values, except for pecan pollen. For example,  $\kappa$  decreased from 0.073±0.006 at 5 °C to 0.057±0.004 at 37 °C for populus tremuloides pollen, and decreased from 0.060±0.001 at 15 °C to 0.050±0.002 at 37 °C for paper mulberry pollen.

385 4 Conclusion and implications

Pollen grains are one of the most abundant types of primary biological aerosol particles in the troposphere and play important roles in many aspects of the Earth system. Hygroscopicity is among the most important physicochemical properties of pollen grains and largely affect their environmental, health and climatic impacts. However, our knowledge in their hygroscopicity is still quite limited, and especially the temperature effect has been rarely explored.

391 In this work we investigated hygroscopic properties of six types of pollen species as a function of RH (up to 95%) at 5 (or 15), 25 and 37 °C. Substantial increase in pollen mass was 392 observed at elevated RH due to water uptake for all the six pollen species. Therefore, change in 393 the mass of pollen grains and their aerodynamic properties at different RH should be taken into 394 account to better understand their transport and deposition in the troposphere. It was found that the 395 mass hygroscopic growth of pollen grains can be well approximated by the modified  $\kappa$ -Köhler 396 equation. The derived  $\kappa$  values at 25 °C ranged from 0.034±0.001 to 0.061±0.007, varying with 397 pollen species. DRIFTS measurements indicated that water adsorption by pollen species were 398 mainly contributed by OH groups of organic compounds contained by pollen grains, and indeed 399 pollen species that contained lower levels of OH groups (relative to C-H groups, as determined 400

401 using transmission FTIR) showed lower hygroscopicity. One exception was corn pollen which contained low levels of OH group but exhibited medium hygroscopicity, suggesting that in 402 addition to chemical composition, other physicochemical properties, such as porosity and internal 403 structure, might also play an important role in determining the hygroscopicity of pollen grains. 404 Due to their moderate hygroscopicity as well as large sizes, pollen grains can thus act as efficient 405 406 giant CCN which may have significant impacts on cloud and precipitation (Johnson, 1982; Feingold et al., 1999; Yin et al., 2000; Posselt and Lohmann, 2008). It is worth noting that only 407 six different pollen species were examined in our work, and hygroscopic properties of other pollen 408 409 species commonly found in the troposphere should be further investigated.

The effect of temperature on the hygroscopicity of pollen grains was systematically 410 investigated in this work. Increase in temperature (from 5 or 15 °C to 25 and 37 °C), a range 411 covering chilling temperature to physiological temperature, led to small but detectable decrease in 412 pollen hygroscopicity. For example,  $\kappa$  values were found to decrease from 0.073±0.006 at 5 °C to 413 0.061±0.007 at 25 °C and to 0.057±0.004 at 37 °C for populus tremuloides pollen, and decrease 414 from 0.060±0.001 at 15 °C to 0.054±0.001 at 25 °C to 0.050±0.002 at 37 °C for paper mulberry 415 pollen. Our measurements at 37 °C (physiological temperature) provide very valuable parameters 416 417 which can be used in numerical models to better understand the transport and deposition of pollen particles in the respiratory system and thus their impacts on human health (Yeh et al., 1996; Broday 418 and Georgopoulos, 2001; Park and Wexler, 2008; Lambert et al., 2011; Longest and Holbrook, 419 420 2012; Tong et al., 2014). Nevertheless, it should be noted that due to the short residence time in the respiratory system, pollen grains and other inhaled particles in general, may not reach 421 422 equilibrium with water vapor in the respiratory tract.

Due to technical challenges, the lowest temperature we could reach in this work was 5 °C, 423 in the range of normal chilling temperatures for vegetative species and also in the expected 424 temperature range at the altitudes of 0.5-2.0 km to which pollen grains can be easily transported. 425 Temperatures in the upper troposphere can be as low as below -70 °C, and it is yet to be explored 426 whether further decrease in temperature to far below 0 °C will lead to large increase in pollen 427 hygroscopicity. As a result, experimental measurements of pollen hygroscopicity at lower 428 temperatures are warranted and would significantly help better understand the transport of pollen 429 grains in the troposphere. Since water vapor has to be adsorbed or condensed on ice nucleating 430 431 particles before heterogeneous ice nucleation can take place (Laaksonen et al., 2016), knowledge in hygroscopicity and water uptake at temperatures below 0 °C would provide fundamental insights 432 into atmospheric ice nucleation, in which pollen grains may play an important role (Pratt et al., 433 2009; Prenni et al., 2009; Hoose et al., 2010; Pöschl et al., 2010; Murray et al., 2012; Creamean et 434 al., 2013; Tang et al., 2018). 435

436

437 Data availability. All the data are available from Mingjin Tang (mingjintang@gig.ac.cn) up on
438 request.

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

440 **Author contribution.** MT, QM and YJL designed the research; WG, CZ, SL and XY did the 441 measurements; MT, QM, YJL and RJH analyzed the results; MT, QM, YJL and RJH wrote the 442 manuscript with contribution from all the co-authors.

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