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# Experimental evidence of a strong image force between highly charged electrosprayed molecular ions and a metal screen

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### 13 Abstract

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We investigated the capturing mechanisms of highly charged macromolecular ions of 15 polyethylene glycol electrosprayed onto a metal screen. Our experiments assessed how the charge 16 state, size of the macromolecular ions, and filtration velocity affected the penetration of the ions 17 through the metal screen. The single fiber efficiencies were plotted as functions of the Peclet 18 number and image force parameter. Highly charged molecular ions had much higher collection 19 efficiencies than neutralized macromolecules, suggesting the presence of a strong image force 20 between the ions and metal surface. The single fiber efficiency by image force was proportional 21 to the square root of an image force parameter predicted by theory. When using the prefactor of 22 9.7 proposed by Alonso et al. (2007), we found fair agreement between the experimental data and 23 theoretical predictions on the collection efficiency of highly charged molecular ions with mobility 24 diameters from 2.6 to 4.8 nm and numbers of electrical charges from 2 to 7. The experimental 25 evidence from our study reveals that image force contributes strongly to the collection of 26 multicharged macromolecular ions by a metal wire screen. 27 28

- 29 Keywords: Image force, Metal screen, Electrospray, Molecular ion
- 30

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## 31 INTRODUCTION

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When a charged particle approaches a neutral solid surface, a short-range force between the 33 particle and induced dipole, a so-called "image force", is created. Image force is a well-known 34 mechanism of particle collection by air filters (Brown, 1993), especially when the particles 35 possess a high number of charges (Yoshioka et al., 1968) or the particle size falls in the 36 37 nanometer range (Alonso et al., 2007; Heim et al., 2010). Yoshioka et al. (1968) measured the collection efficiency of electrically charged oil droplets with a 1 µm diameter by a neutral glass 38 fiber filter. The number of charges per droplet was controlled using a pin-to-plate type discharge 39 electrode. The collection efficiency of 60-charged particles (1 µm) was about 40 % higher than 40 that of uncharged particles in their experiment. Lundgen and Whitby (1965) measured the 41 collection efficiency of highly charged (-300 to +320 elemental charges) particles with 0.1 and 1 42 43 um diameters using electrically neutral filters made of felt, urethane, and glass fiber. They reported that the collection efficiency increased from 16 % (uncharged particles) to over 99 % 44 45 when the particles carried +320 electrical charges. To evaluate the effect of image force in a diffusion limited regime (image force parameter,  $K_{IM} < 10^{-5}$ ), Alonso *et al.* (2007) experimentally 46 measured the collection efficiency of nanoparticles (ZnCl<sub>2</sub> and NaCl) with diameters from 25 to 47 48 65 nm by a metal screen (aluminum and stainless steel). The image force had contributed significantly to the mechanism by which the charged nanoparticles, even the particles with small 49 numbers of electrical charges, were collected in their experiment. 50

As reported in previous studies, image force has a significant influence on the mechanism by which highly charged nanoparticles are collected. Few reports, however, have investigated the effect of image force on the collection of very small particles (< 10 nm). New factor to consider emerge in a size regime of < 10 nm, such as the thermal rebound (Wang and Kasper, 1991) and the effects of Brownian diffusion and image force in combination (Alonso *et al.*, 2007). Heim *et al.* (2010) investigated the collection efficiencies of tungsten oxides nanoparticles (1.2 to 8 nm, neutral and monovalent) and electrosprayed molecular ions by metal meshes (nickel and stainless steel). The efficiencies of nanoparticle collection in their experiments were in good agreement with the predicted collection efficiency of particles with diameters larger than 3 nm. They also found, however, that image force has only a minor effect on the collection efficiency of nanoparticles, because the number of charges of their test particles was unity.

We studied the effect of image force on the particle collection in the course of investigation of 62 thermal rebound of sub-10 nm particles. Since we have neither definite generation method nor the 63 measurement methods for sub-10 nm particles, we employed macromolecular ions as a 64 monodisperse test particles generated by an electrospray, which is a powerful tool for producing 65 molecular ions (Yamashita et al., 1984). However, we found that the electrospray generates 66 highly charged macromolecular ions so that the collection efficiency of these ions was 67 significantly affected by the image force. Consequently, we must account for the image force in 68 the interpretation of collection efficiency data for elucidating the thermal rebound of sub-10 nm 69 70 particles.

In the present study, we generated highly charged polyethylene glycol (PEG) molecular ions (from +2 to +7) with diameters below 5 nm by an electrospraying PEG with various molecular weights. We measured the collection efficiency of highly charged molecular ions through wire screens as a function of mobility diameters and numbers of charges and compared the experimental data with the empirical equation reported by Alonso *et al.* (2007) in order to examine whether the empirical equation obtained for particles larger than 10 nm holds for sub-10 nm particles.

#### **COLLECTION OF CHARGED NANOPARTICLES BY IMAGE FORCE**

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81 The influences of image force were evaluated by a dimensionless image force parameter,  $K_{\text{IM}}$ , 82 given by

83

84 
$$K_{\rm IM} = \left(\frac{\varepsilon_{\rm f} - 1}{\varepsilon_{\rm f} + 2}\right) \frac{Cq^2 e^2}{12\pi^2 \mu u \varepsilon_0 d_{\rm p} d_{\rm f}^2}$$
(1)

85

where,  $\varepsilon_{\rm f}$  and  $\varepsilon_0$  are dielectric constants of the fiber and vacuum, *C* is the Cunningham slip correction factor, *q* is the number of charges on the particle, *e* is the elementary charge,  $\mu$  is the air viscosity, *u* is the air flow velocity, *d*<sub>P</sub> is the particle diameter, *d*<sub>f</sub> is the fiber diameter. Single fiber efficiency by image force is known to be proportional to the square root of *K*<sub>IM</sub> such that,

90

91 
$$\eta_{\rm IM} = \alpha \sqrt{K_{\rm IM}}$$
 (2)

92

where,  $\alpha$  is a constant of proportionality, which takes different value in different reports. When 93 Yoshioka et al. (1968) measured the collection efficiency of dioctyl phthalate by a glass filter mat, 94 they reported an  $\alpha$  value of 2.3 from experiments using 1 mm particles ( $10^{-6} < K_{IM} < 10^{-3}$ ). A few 95 years earlier, Lundgen and Whitby (1965) reported an  $\alpha$  value of 1.5 from experiments on solid 96 spherical aerosol particles with diameter ranging from 0.1  $\mu$ m to 1  $\mu$ m (2×10<sup>-6</sup> <  $K_{IM}$  < 3×10<sup>-2</sup>). 97 In the diffusion regime, the total single fiber efficiency,  $\eta_{\rm T}$ , is expressed as the sum of the single 98 fiber efficiencies of diffusion, image force, and the combined effect of diffusion and image force 99 100 as,

102 
$$\eta_{\rm T} = \eta_{\rm D} + \eta_{\rm IM} + \eta_{\rm DIM} \tag{3}$$

104 The single fiber collection efficiency of diffusion,  $\eta_D$  is expressed by the following equation 105 (Cheng and Yeh, 1980):

106

107 
$$\eta_{\rm D} = 2.7 P e^{-2/3}$$
 (4)

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109 where, *Pe* is the Peclet number given by

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111 
$$Pe = \frac{ud_{\rm f}}{D}$$
(5)

112

where *D* is the diffusion coefficient. The third term of Eq. (3),  $\eta_{\text{DIM}}$ , is the single fiber efficiency by a combination of two capturing mechanisms, diffusion and image force.

Alonso *et al.* (2007) found that the pure image force effect was negligible in the diffusion regime (200 < Pe < 3600,  $10^{-7} < K_{IM} < 10^{-5}$ ) and obtained the following equation for  $\eta_{DIM}$ (combined effect):

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119 
$$\eta_{\rm DIM} = 29.7 K_{\rm IM}^{0.59}$$
 (6)

120

121 They also fitted the experimental data by assuming that  $\eta_{\text{DIM}}$  was proportional to the square 122 root of the  $K_{\text{IM}}$ , and it also shows reasonable agreement between the experimental data when  $\alpha =$ 123 9.7 as follows:

125 
$$\eta_{\text{DIM}} = 9.7\sqrt{K_{\text{IM}}}$$

Fig. 1 shows the size dependency of single fiber efficiency by image force,  $\eta_{IM}$ , calculated by Eq. (7) with changing numbers of charges, q ( $u = 0.25 \text{ m s}^{-1}$ ). The solid line and dashed lines respectively represent the single fiber efficiency by pure diffusion and by diffusion and image force combined. As the figure shows, the single fiber efficiency increases with q and decreases with the particle diameter. The image force effect is clearly significant when the particles carry more than several electrical charges. Highly charged nanoparticles of this type can be generated by electrospray atomization.

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#### 135 METHODS

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#### 137 Preparation of multicharged PEG molecular ions

Fig. 2 shows the experimental setup used in this study. Highly charged electrosprayed 138 139 molecular ions were used as test particles in the filtration experiments. Polyethylene glycol (PEG) with the monodispersed molecular weight from 2,000 to 21,300 were purchased from Sigma 140 Aldrich ( $M_W = 10000$ ), Wako ( $M_W = 2000$ , 4600) and Agilent Technologies ( $M_W = 21300$ ). PEG 141 (0.005 to 0.043 wt%) and 0.1 wt% ammonium acetate were dissolved in a water/methanol (50:50) 142 solution for the electrospray atomization. The mixture of water/methanol solution, and the 143 addition of ammonium acetate have been widely employed in order to increase the electrical 144 conductivity of precursor solution in electrospray process (Lenggoro et al., 2002). In addition, the 145 correlations between PEG concentration and charging state were reported in our previous 146 research (Maekawa et al., 2014). 147

The PEG solution was introduced to the capillary (Hamilton, Model 7747-02, inner diameter of 148 110 μm, outer diameter of 240 μm) via a syringe pump at a constant flow rate of 100 to 200 μL 149  $h^{-1}$ . High voltage (2.5 to 3 kV) was applied to the capillary to obtain a stable cone jet under 150 observation by a CCD camera (Sony, Model XC-75). as shown in Fig. 3. The gap distance 151 between syringe tip to electrical ground was set to 0.3 cm. The generated droplets were carried by 152 dry clean air at flow rate of 3 to 5 L min<sup>-1</sup>. Part of the gas flow was introduced into a differential 153 mobility analyzer (DMA) (Laboratory-made, Vienna type, Winklmayr et al. 1991) to obtain 154 mono-mobility particles. The number concentrations of the DMA classified particles were 155 monitored using a condensation particle counter (CPC) (TSI, Model 3776, Fig. 2(a)). 156

The charging state of the test particles was investigated by introducing the DMA-classified PEG molecular ions to a neutralizer ( $^{241}$ Am) and measuring the mobility distribution of the neutralized PEG molecules measured by a Nano-SMPS (TSI, Model 3085 nano-DMA and the CPC, Fig. 2(b)). At the test particles were very small (< 5 nm), the molecular ions classified by the Nano-SMPS were assumed to be monovalent. The number of electrical charges of the PEG ions could therefore be estimated as a function of the diameter by changing the 1<sup>st</sup> DMA voltage (for mobility scanning) and measuring the size distributions of the ions with the Nano-SMPS.

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## 165 Evaluation of collection efficiency

Mobility-classified molecular ions of known sizes and numbers of charges were introduced to the filtration experiment system. Fig. 4 shows the experimental setup for measuring the filtration efficiency. The effect of the image force in the diffusion regime was assessed by comparing the penetration data of highly charged PEG ions (Fig. 4(a)) with the penetration data of neutralized PEG molecules (Fig. 4(b)). The diffusional loss of the ions during transport was cancelled by measuring the penetration using two identical filter holders (Alonso *et al.*, 2007; Heim *et al.*, 2010) arranged shown in Fig. 4. Most of the tubes and connections were made of metal
(electrically grounded) and set at the shortest possible lengths to minimize the diffusional loss of
ions during transport.

A stainless steel (SUS) screen (fiber diameter,  $d_f = 30 \ \mu m$ , packing density,  $\alpha = 0.177$ ) was installed in one of the filter holders for use as the model filter. The diameter of the effective filtration area was 24 mm. The filtration velocity was adjusted from 0.25 to 0.40 m s<sup>-1</sup> by adding clean makeup air before the ions entered the filter holders. The flow was switched between the filter and the blank holder by three-way valves. The number concentration with and without filtration was measured by the CPC.

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## 182 **RESULTS AND DISCUSSION**

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#### 184 Characterization of the charged molecular ions

Fig. 5 shows contour plots of the electrical mobility (multicharged particles measured by the 1<sup>st</sup> 185 DMA) against the particle diameter (singly charged particles measured by the 2<sup>nd</sup> DMA) for (a) 186 PEG2000, (b) PEG4600, (c) PEG10000, and (d) PEG21300. The number concentration is 187 represented as a difference of color (red is the highest concentration). Each of upper panels shows 188 the CPC count plotted against particle diameters measured by directly introducing electrosprayed 189 ions into the Nano-SMPS with a neutralizer. Two kinds of products were identified in the size 190 distributions. The first peak observed at the smaller particle sizes could be attributed to single 191 molecular ions, since the diameter of these first peaks increased with the increasing molecular 192 193 weight of the PEG. The second peak was thought to be composed of residual nanoparticles generated by the drying of the electrosprayed droplets containing multiple PEG molecules. The 194 diameter of this peak increased with increasing concentrations of the PEG solutions. 195

Only the molecular ions from the first peaks were used as test particles for the filtration experiment, as the particles in the second peak may have been mixtures of multicharged residual particles of various sizes and numbers of charges. Each of the solid lines in Fig. 5 represents the relationship between the electrical mobility of the particles,  $Z_p$ , and the particle diameter,  $d_p$ , as a function of the number of charges, q, given by the following Eqs. (8) and (9):

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202 
$$Z_{\rm p} = \frac{qeC(d_{\rm p})}{3\pi\mu d_{\rm p}}$$
(8)

203

204 
$$C = 1 + \frac{2\lambda}{d_{\rm p}} \left[ 1.246 + 0.420 \exp\left(-0.87 \frac{d_{\rm p}}{2\lambda}\right) \right]$$
 (9)

205

where *C* is the slip correction factor, and  $\lambda$  is the mean free path.

By comparing the lines with contour plots we find, that the PEG ions have 2 to 7 charges, 207 208 depending on the molecular weight. Table 1 summarizes the relationship of the molecular weight with the mobility diameter and number of electrical charges. Disappointingly, the resolution of 209 the 1st DMA used for mobility classification was too low to accurately classify molecular ions 210 with the different numbers of charges (Saucy et al. 2004). Its resolution was sufficient, however, 211 to eliminate residual nanoparticles, as shown in Fig. 5. We therefore compared our experimental 212 data with the theoretical values by assuming that all of the test particles had the same number of 213 charges (see Table 1). 214

215

#### 216 Collection of highly charged molecular ions

Fig. 6 shows penetration data of the four PEG species through an SUS wire screen with and without electrical neutralization at filtration velocities of  $0.25 \text{ m s}^{-1}$ . The solid line and dashed 219 line represent the theoretical penetration curves for uncharged and singly charged macromolecules, respectively. As the figure shows, experimentally obtained penetration of 220 221 neutralized macromolecules (solid circles) decreases with decreasing diameter, which suggests that the main capturing mechanism is diffusion. It also shows fair agreement with the predicted 222 penetration data for uncharged macromolecules (only diffusion) or singly charged ions. When the 223 charged particles are neutralized to the equilibrium charging state, the fractions of the charged 224 particles are estimated to be about 1% for 2 nm and 3% for 5 nm particles by Fuchs' charging 225 theory (Fuchs, 1963). From these estimates, conclude that the experimentally obtained 226 penetration values should closely agree with the predicted curve for uncharged macromolecules. 227 We find, however, that the predicted curve diverges from the experimentally obtained values, 228 possibly due to the residual electrical charge on the particles. On the other hand, we clearly see 229 that the multicharged ions (open circles) have a much lower penetration than the neutralized 230 species. We attribute this result to the significant contribution of the image force between the 231 multicharged particles and SUS wire screen. 232

To analyze the effect of image force, we converted the penetration, P, to the single fiber efficiency,  $\eta$ , given by Eq. (10):

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236 
$$P = \exp\left(-\frac{4}{\pi}\frac{\alpha}{1-\alpha}\frac{L}{d_{\rm f}}n_{\rm screen}\eta\right)$$
(10)

237

where  $\alpha$  is the packing density of the wire screen, *L* is thickness of wire screen, and *n*<sub>screen</sub> is number of wire screen. Fig. 7 shows the experimental single fiber efficiencies of neutralized macromolecules and multiply charged ions obtained from Eq. (10), plotted as functions of the Peclet number, *Pe* (Eq. (5)). Each symbol in Fig. 7 represents the series of data for the specific

molecular weights of PEG (shown in Table 1) with changing filtration velocity. The experimental 242 single fiber efficiency of the multicharged species (open symbols) showed various tendencies at 243 different molecular weights (diameters) and charge states. In contrast, the single fiber efficiencies 244 of the neutralized macromolecules determined experimentally (closed symbols) could almost be 245 summarized into a straight line as a function of Pe and agreed fairly well with the predicted 246 efficiencies for both neutralized (solid line) and singly charged ions (broken line), as previously 247 mentioned. The broken, dashed, and dotted lines in Fig. 7 represent the theoretical single fiber 248 efficiencies for multicharged ions (1 to 11 charges) obtained from the sum of Eqs. (4) and (7). As 249 previously mentioned (Fig. 5), the numbers of charges obtained experimentally were 4.4 for 250 PEG<sub>4600</sub>, 6.4 for PEG<sub>1000</sub>, and 7.2 for PEG<sub>21300</sub>. As the figure illustrates, the experimental single 251 252 fiber efficiencies of multicharged ions increased with increasing number of charges and agreed fairly well with the predicted lines with corresponding numbers of charges. This result provides 253 quantitative evidence of a strong image force between the multicharged molecular ions and the 254 SUS wire screen even in the diffusion regime. 255

To reiterate, Alonso *et al.* (2007) reported that a prefactor  $\alpha$  of 9.7 validly applied to estimation 256 257 of the image force for multicharged (maximum 3) nanoparticles in the diffusion regime (mobility diameters from 25 to 60 nm). In the present study, we extend the validation of the prefactor down 258 to molecular ions as small as 2.6 nm (Fig. 6). In contrast, Heim et al. (2010) performed filtration 259 260 experiments for singly charged nanoparticles in the diameter range from 1.2 to 8 nm using a metal screen. The dominance of Brownian diffusion as the mechanism of particle capture in their 261 experiment reduced the contribution of pure image force to a negligible level. They tentatively 262 263 explained this dominant effect of Brownian diffusion by the weak image force it for small numbers of charges (single), a phenomenon identified in Fig. 7 (the small difference between 264 neutral and singly charged particles). 265

In summary, Fig. 8 plots the single fiber efficiency for the charged particles (the sum of pure 266 image force plus image force and diffusion in combination),  $\eta_{IM} (= \eta_T - \eta_D)$ , as a function of the 267 image force parameter KIM. Our experimental data and Alonso's results are plotted as open 268 symbols, and closed symbols, respectively. The solid line represents the single fiber efficiency 269 predicted by Eq. (7). As the figure shows, our experimental results agree fairly well with the line 270 predicted by Alonso's prefactor of 9.7. Our study confirms that Alonso's prefactor of 9.7 can be 271 applied in our experimental ranges ( $d_p = 2.6 \sim 4.8$  nm,  $q = 2 \sim 7$ ), which correspond to  $10^{-3} < K_{IM}$ 272  $< 10^{-2}$ . 273

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## 275 CONCLUSIONS

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We prepared multicharged ions carrying 2 to 7 elementary charges by an electrospray method, 277 and investigate the mechanism by which they were collected by a SUS wire screen. The 278 multicharged molecular ions had lower penetration than the neutralized molecules due to the 279 image force acting between the multicharged molecular ions and SUS wire screen. The single 280 fiber efficiency by image force also turned out to be proportional to the square root of an image 281 282 force parameter predicted by theory, and the experimental and theoretical values showed fair agreement when we using the prefactor of 9.7 proposed by Alonso et al., even in the KIM ranges 283 from  $10^{-3}$  to  $10^{-2}$ . This study demonstrates the strong contribution of image force in the collection 284 of multicharged molecular ions by an SUS wire screen. 285

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## 287 **REFERENCES**

- 289 Alonso, M., Alguacil, F.J., Santos, J.P., Jidenko, N. and Borra, J.P. (2007). Deposition of ultrafine aerosol particles on wire screens by simultaneous diffusion and image force. J. Aerosol Sci. 38: 290 291 1230-1239.
- Brown, R.C. (1993). Air filtration: An integrated approach to the theory and applications if 292 293 fibrous filters, Pergamon Press, Oxford.
- Cheng, Y.S. and Yeh, H.C. (1980). Theory of a screen-type diffusion battery. J. Aerosol Sci. 11: 294 313-320.
- Fuchs, N.A. (1963). On the stationary charge distribution on aerosol particles in a bipolar ionic 296 atmosphere. Geofis. Pura. Appl. 56: 185-193. 297
- Heim, M., Attoui, M. and Kasper, G. (2010). The efficiency of diffusional particle collection onto 298
- 299 wire grids in the mobility equivalent size range of 1.2-8 nm. J. Aerosol Sci. 41: 207-222.
- Lundgren, D.B. and Whitby, K.T. (1965). Effect of particle electrostatic charge on filtration by 300 fibrous filters. Ind. Eng. Chem. Process Des. Dev. 4: 345-349. 301
- Lenggoro, I. W., Xia, B., Okuyama, K., and De la Mora, J. F. (2002). Sizing of colloidal 302 nanoparticles by electrospray and differential mobility analyzer methods. Langmuir, 18: 4584-303 4591. 304
- Maekawa, T., Tokumi, T., Higashi, H., Seto, T. and Otani, Y. (2014). Effect of solution 305 concentration on breakup of electro-sprayed droplets and emission of solute ions. Kagaku 306 Kogaku, 40: 5-11. 307
- Saucy, D.A., Ude, S., Lenggoro, I.W. and De la Mora, J.F. (2004). Mass analysis of water-soluble 308
- polymers by mobility measurement of charge-reduced ions generated by electrosprays. Anal. 309
- 310 Chem. 76: 1045-1053.

Wang, H.C. and Kasper, G. (1991). Filtration efficiency of nanometer-size aerosol particles. J. 311 Aerosol Sci. 1: 31-41. 312

- Winklmayr, W., Reischl, G.P., Lindner, A.O. and Berner, A. (1991). A new electromobility spectrometer for the measurement of aerosol size distributions in the size range from 1 to 1000 nm. *J. Aerosol Sci.* 22: 289-296.
- 316 Yamashita, M. and Fenn, J.B. (1984). Electrospray ion source. Another variation on the free-jet
- 317 theme. J. Phys. Chem. 88: 4451-4459.
- 318 Yoshioka, N., Emi, H., Hattori, M. and Tamori, I. (1968). Effect of electrostatic force in the
- 319 filtration efficiency of aerosols. *Kagaku Kogaku* 32: 815-820.
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## **Table Titles**

**Table 1.** Mobility diameters and charging characteristics of PEG ions.

Molecular weight of PEG [g/mol]	Concentration of PEG [wt%]	Mobility diameter [nm]	Number of charges [-]	Average number of charges [-]
2,000	0.005	2.6	1~2	2.25
4,600	0.023	3.1	2~7	4.38
10,000	0.02	4.1	3~9	6.37
21,300	0.043	4.8	5~12	7.27

 Table 1. Mobility diameters and charging characteristics of PEG ions.

325	Figure Captions
326	Fig. 1. Single fiber collection efficiency by diffusion and by image force.
327	Fig. 2. Experimental setup to prepare the PEG ions.
328	Fig. 3. Microscopic image of the cone-jet.
329	Fig. 4. Experimental setup to measure the collection efficiency.
330	Fig. 5. CPC count and tandem result of PEG ions.
331	Fig. 6. Penetration of PEG ions as a function of mobility diameter.
332	Fig. 7. Single fiber collection efficiencies of PEG ions as functions of the Peclet number.
333	Fig. 8. Single fiber efficiency by image force with Alonso's prefactor.
334	



Fig. 1.





Fig. 2.



Fig. 3









Fig. 5.



Fig. 6.



Fig. 7.



Fig. 8.