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Title	Effects of decreasing activated carbon particle diameter from 30 mu m to 140 nm on equilibrium adsorption capacity
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#### **Research Highlights**

- Micromilling produced carbon particles with D50s as low as 140 nm.
- Adsorption capacity decreased as D50 decreased from a few micrometers to 140 nm.
- The adsorption capacity decrease was due to carbon oxidization during micromilling.
- Inhibition of carbon oxidation attenuated the decrease in adsorption capacity.
- Adsorption capacity was negatively correlated with particle oxygen content.



1	Effects of Decreasing Activated Carbon Particle Diameter from 30 µm to 140 nm on
2	Equilibrium Adsorption Capacity
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#### 13 Abstract

14 The capacity of activated carbon particles with median diameters (D50s) of  $>\sim 1 \mu m$  for adsorption of hydrophobic micropollutants such as 2-methylisolborneol (MIB) increases 15 with decreasing particle size because the pollutants are adsorbed mostly on the exterior 16 (shell) of the particles owing to the limited diffusion penetration depth. However, particles 17 with D50s of <1 µm have not been thoroughly investigated. Here, we prepared particles 18 19 with D50s of ~30 µm to ~140 nm and evaluated their adsorption capacities for MIB and several other environmentally relevant adsorbates. The adsorption capacities for low-20 molecular-weight adsorbates, including MIB, deceased with decreasing particle size for 21 D50s of less than a few micrometers, whereas adsorption capacities increased with 22 decreasing particle size for larger particles. The oxygen content of the particles increased 23 24 substantially with decreasing particle size for D50s of less than a few micrometers, and oxygen content was negatively correlated with adsorption capacity. The decrease in 25 adsorption capacity with decreasing particle size for the smaller particles was due to particle 26 27 oxidation during the micromilling procedure used to decrease D50 to ~140 nm. When oxidation was partially inhibited, the MIB adsorption capacity decrease was attenuated. For 28 high-molecular-weight adsorbates, adsorption capacity increased with decreasing particle 29 size over the entire range of tested particle sizes, even though particle oxygen content 30 increased with decreasing particle size. 31

32

#### 33 Keywords

34 MIB, micro-grinding, bead mill, oxygen content, oxidation

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#### 37 **1. Introduction**

38 Superfine powdered activated carbon (SPAC), which is produced by micromilling of normal powdered activated carbon (PAC), is a new option for efficiently removing natural 39 organic matter, disinfection byproduct precursors, organic micropollutants, and taste and 40 odor compounds from water (Bonvin et al. 2016, Dunn and Knappe 2013, Heijman et al. 41 2009, Matsui et al. 2007, Matsui et al. 2004, Matsui et al. 2005, Wang et al. 2011). SPAC 42 offers many advantages over PAC, such as higher adsorption capacity (Ando et al. 2010, 43 44 Ando et al. 2011, Bonvin et al. 2016, Matsui et al. 2010), faster adsorption kinetics (Dudley 2012, Matsui et al. 2009a, Matsui et al. 2011, Wang et al. 2011), and the ability to attenuate 45 the buildup of transmembrane pressure when used before microfiltration (Ellerie et al. 2013, 46 Li 2014, Matsui et al. 2007, Matsui et al. 2009b). 47

The higher adsorption capacity of SPAC is of both practical and theoretical interest because in previous studies it had been believed that adsorption occurred in the internal pores of activated carbon (AC) particles and that equilibrium adsorption capacity was therefore independent of particle size (Letterman et al. 1974, Najm et al. 1990, Peel and Benedek 1980).

Previous investigations have reported that the change in the AC particle size caused by micro-milling does not result in any substantial change in the internal pore area (Ando et al. 2010, Matsui et al. 2014, Pan et al. 2017). Therefore, the higher adsorption capacity of SPAC is not related to the internal pore area. More recently, the higher adsorption capacity has been explained in terms of the shell adsorption model (SAM), which holds that adsorbates are adsorbed mainly on the exterior (shell) of particles owing to the limited diffusion penetration depth (Ando et al. 2011, Matsui et al. 2011, Matsui et al. 2013, Matsui

et al. 2014). The decrease in particle size upon micromilling increases the exterior surface 60 area, thus enhancing adsorption capacity, as long as the particle radius is larger than the 61 diffusion penetration depth, which depends on the adsorbate and the AC. For example, it 62 has been shown that 2-methylisoborneol (MIB), a taste and odor compound that is a target 63 of water treatment processes, penetrates to a depth of 4-6 µm in some coconut-shell-based 64 and wood-based PACs and that SPACs (particle radius,  $0.5-1 \mu m$ ) produced from the PACs 65 have higher MIB adsorption capacities than the PACs (particle radius, 5-10 µm) (Matsui 66 67 et al. 2015, Matsui et al. 2014).

68 The SAM also suggests that if the adsorbate penetration depth is a few micrometers, adsorption capacity may not increase when the AC particle radius is decreased from a few 69 micrometers to less than a micrometer. However, adsorption capacities and characteristics 70 of ACs in this particle size range have not been thoroughly investigated. Micromilling is a 71 new nanotechnology that has enabled production of particles with median diameters (D50s) 72 from the submicrometer range down to hundreds of nanometers, and even smaller (Mende 73 et al. 2003, 2004). This technology has been applied to materials such as corundum (Stenger 74 et al. 2005), aluminum nitride (Jia et al. 2015, Qiu et al. 2006), Li<sub>4</sub>Ti<sub>5</sub>O<sub>12</sub> (Han et al. 2012), 75 barium sulfate (Patel et al. 2014), silica (Patel et al. 2015), and crystalline pyrene (Flach et 76 al. 2016). Our research group (Pan et al. 2015, Pan et al. 2017) and others (Amaral et al. 77 2016, Partlan et al. 2016) have produced AC particles with D50s of 200-250 nm. 78

Several full-scale SPAC-microfiltration plants are already being used for drinking
water production (Kanaya et al. 2015); in these plants, SPAC with D50s of 1–2 μm is
produced onsite and added before membrane filtration. The use of smaller particles (e.g.,
150-nm diameter) might enhance adsorptive removal efficiency.

- In this study, we produced ACs with D50s of 30 µm to 140 nm and evaluated the effects
  of particle size on their adsorption capacities for various adsorbates.
- 85 2. Experimental

86 2.1 ACs

87	Five commercially available PACs (Carbons A–E, Table 1) were milled normally (see
88	sec. 2.2.1) to produce SPACs and submicrometer SPACs (SSPACs). The PACs, SPACs,
89	and SSPACs were given unique four-part designations specifying particle size, parent PAC
90	(A–E), production method, and batch number (where applicable). For the first term, PAC
91	designates particles with D50s of 12.0–28.0 $\mu$ m; SPAC <sub>L</sub> , D50s of 3.80–8.10 $\mu$ m; SPAC <sub>s</sub> ,
92	D50s of 0.794–1.80 µm; and SSPAC, D50s of 0.137–0.230 µm. The third term (used only
93	for SPAC <sub>L</sub> , SPAC <sub>S</sub> , and SSPAC) indicates the production method: "N" for normal milling,
94	"NET" for extended-time normal milling, "OI" for oxidation-inhibiting milling, and "R" for
95	rinsing with pure water (see sec. 2.2 and Tables S1 and S2). For example, SSPAC-C-N-2
96	is the second batch of an SSPAC produced from PAC-C under normal milling conditions.

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98 2.2 SPAC and SSPAC production

99 SPAC<sub>L</sub> was produced with a closed-chamber ball mill (Nikkato, Osaka, Japan) with 5-100 and 10-mm-diameter balls. SPAC<sub>s</sub> and SSPAC were produced with a bead mill with a re-101 circulation system (LMZ015, Ashizawa Finetech, Chiba, Japan) with (0.1- or 0.3-mm-102 diameter  $ZrO_2$  beads. The next two sections describe the milling procedures.

103

104 2.2.1 Normal milling

As-received PACs (designated as, e.g., PAC-A) were slurried in pure water (Milli-Q Advantage, Millipore, Bedford, MA, USA) at a concentration of ~15% (w/w) and then milled in the ball mill at <45 rpm for 4–5 h to afford particles with D50s of 3.03–3.90  $\mu$ m (designated as, e.g., SPAC<sub>L</sub>-A-N). The slurry was collected from the chamber of the ball mill with pure water and then diluted to ~1% (w/w).

The diluted AC slurry (400 mL) was then milled with the bead mill with 0.3-mmdiameter beads at a rotational speed of 2590 rpm in recirculation mode for 20–30 min to afford particles with D50s of <1.47  $\mu$ m but >0.915  $\mu$ m (designated as, e.g., SPACs-A-N). AC particles with D50s of 137–198 nm were produced from SPAC<sub>L</sub> or SPAC<sub>S</sub> using the bead mill with 0.1-mm-diameter beads at 3884 rpm in re-circulation mode for 1.5–2 h. In additional experiments, the milling time was extended to 5–7 h. During milling, the slurry was kept at <28 °C by means of a cooling system.

117

118 2.2.2 Oxidation-inhibiting milling

As-received PACs were dried in a vacuum chamber for at least 1 h, and then the chamber was filled with N<sub>2</sub> at atmospheric pressure. The PACs in the N<sub>2</sub>-filled chamber were rinsed with pure water three times. The PACs obtained in this way are designated as, e.g., PAC-C-R.

The rinsed PAC slurries were adjusted to a concentration of ~15% (w/w) with pure water, transferred to the ball mill filled with N<sub>2</sub> gas, and milled under the conditions used for normal milling. Because collecting the AC slurries under N<sub>2</sub> after milling was difficult, they were collected in air, but as quickly as possible. The collected slurries were adjusted to a concentration of ~1% (w/w) with pure water. The dissolved oxygen concentration was then reduced to <1 mg/L by allowing N<sub>2</sub> gas to diffuse into the slurries for >1 h. The slurries obtained in this way are designated as, e.g., SPAC<sub>L</sub>-C-OI.

SPACs-OI was produced by bead milling of SPAC<sub>L</sub>-OI under conditions in which the slurry was purged with N<sub>2</sub> (Fig. 1); otherwise, the milling conditions were the same as those used for normal bead milling. After milling, the slurries were kept under vacuum for >1 h and then placed under N<sub>2</sub> at atmospheric pressure. The slurries obtained in this way are designated as, e.g., SPAC<sub>S</sub>-C-OI.

SSPAC-OI was produced by bead milling SPAC<sub>L</sub>-OI (400–500 mL) under N<sub>2</sub> purging,
similar to those used to produce SPAC<sub>s</sub>, for 3–4 h at 2590 rpm with 0.1- or 0.3-mmdiameter beads.

All the AC slurries were stored at 4 °C after vacuum conditioning to remove air from
the pores.

140

141 2.3 AC characterization

Particle size distributions of all the ACs were measured by laser-light scattering
(Microtrac MT3300EXII, Nikkiso, Tokyo, Japan) in transmission mode. Samples of stock
slurries were pretreated with a dispersant (Triton X-100, Kanto Chemical, Tokyo, Japan;
final concentration, 0.08% w/v) and ultrasonically dispersed (150 W, 19.5 kHz, 4 min for
50 mL AC slurry) before measurement.

SSPAC particles were observed by field emission scanning electron microscopy (JSM7400F, JEOL, Tokyo, Japan) at an accelerating voltage of 5.0 kV and a working distance
of 3.0 mm. The particles were not precoated, because the thickness of the coating layer
might have hindered accurate observation. SPACL, SPACs, and PAC particles were not

observed microscopically, because images of these particles had already been obtained
(Ando et al. 2010) and because the particles sizes were visually confirmed.

The oxygen, hydrogen, nitrogen, and sulfur contents of the ACs were determined by 153 elemental analysis (Vario EL Cube, Elementar Japan, Yokohama, Japan) with thermal 154 conductivity and infrared detection. For this purpose, the AC slurries were dried by a 155 vacuum cooling method to prevent oxidation during drying. Before elemental analysis, the 156 dried samples were placed in small tin capsules and weighed out. The capsules containing 157 sample were stored in an acrylic container under a vacuum. After atmospheric pressure N2 158 was introduced to the box, the capsules were promptly transferred to the elemental analyzer. 159 The oxygen content was measured in O mode, and the other elements were measured in 160 CHNS mode. Ash contents were determined by igniting oven-dried AC samples in a muffle 161 furnace at 550 °C. Averages of triplicate measurements are reported for each AC. 162

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#### 164 2.4 Adsorbates and working solution

MIB was the main target compound in this study. MIB concentrations were determined 165 in two ways. Either the m/z 95 peak of MIB was detected with a purge-and-trap concentrator 166 (Aqua PT 5000 J, GL Sciences, Tokyo, Japan) coupled to a gas chromatograph-mass 167 spectrometer (GCMS-QP2010 Plus, Shimadzu, Kyoto, Japan) using geosmin-d<sub>3</sub> (m/z 115) 168 as an internal standard. Or the m/z 108 peak of MIB was detected by headspace solid-phase 169 microextraction (PAL RSI 85, Agilent Technologies Japan, Ltd., Tokyo, Japan) coupled to 170 a gas chromatograph-mass spectrometer (7820A/5977 E MSD, Agilent) using 2,4,6-171 trichloroanisole- $d_3$  (m/z 195) as an internal standard. 172

On the basis of environmental relevancy, molecular size, and hydrophobicity, nine other adsorbates were tested in addition to MIB: geosmin (log Kow, 3.57; 182 Da),

dimethylphosphorothidate-oxon (DMTP-oxon; log Kow, 0.77; 286 Da), benzothiazole (log 175 Kow, 2.09; 135 Da), phenol (log Kow, 1.46; 94 Da), para-nitrophenol (log Kow, 1.91; 139 176 Da), salicylic acid (log Kow, 2.06; 138 Da), poly(styrenesulfonic acid sodium salt) with an 177 average molecular weight (MW) of 210 Da (PSS-210), PSS-1100 (average MW, 1100 Da), 178 and SNOM (Suwannee River natural organic matter, International Humic Substance 179 Society). Note that the log Kow values listed above are from EPI Suite<sup>™</sup> database (ver. 4, 180 181 U.S. Environmental Protection Agency). Geosmin concentrations were determined by a method similar to that used for MIB; m/z 112 was attributed to geosmin. DMTP-oxon was 182 183 quantified by hybrid quadrupole-orbitrap mass spectrometry (Q Exactive, Thermo Fisher Scientific, Waltham, MA, USA) combined with liquid chromatography (UltiMate3000 LC 184 systems, Thermo Fischer Scientific) using mepronil as an internal standard (DMTP-oxon, 185 186 m/z 287; mepronil, m/z 270). The other adsorbate concentrations were measured by UVvis spectrophotometry (UV-1800, Shimadzu, Kyoto, Japan) at 269.5, 252, 296, 317, and 187 261 nm for phenol, benzothiazole, salicylic acid, para-nitrophenol, and PSS-210/PSS-1100, 188 respectively. 189

Except for PSS-210 and PSS-1100 (Sigma-Aldrich, St. Louis, MO, USA), all chemical reagents were purchased from Wako Pure Chemical Industries (Osaka, Japan). The working solution was organic-free water with the ion composition used in previous research (Table S3) (Matsui et al. 2015, Pan et al. 2016). The working solutions were adjusted to pH 7.0  $\pm$ 0.1 with HCl or NaOH as required. The solution pH was maintained constant during adsorption tests probably due to bicarbonate buffer.

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197 2.5 Batch adsorption tests

For each target adsorbate, an aliquot (100 or 110 mL) of working solution containing the adsorbate was transferred to a 110-mL vial; AC was immediately added; and the vial

was manually shaken and then agitated on a mechanical shaker for 1 week or 3 days at 200 20 °C in the dark. Preliminary experiments confirmed that adsorption equilibrium was 201 reached within 1 week for the MIB-PAC combination (Matsui et al. 2013, Matsui et al. 202 2012), 3 days for MIB-SPAC/SSPAC, and 1 week for the other 8 adsorbates (for all the 203 ACs). Control tests of blank solutions containing adsorbates but no AC confirmed that 204 changes in adsorbate concentration during long-term mixing were negligible. After shaking, 205 206 water samples were filtered through a 0.2-µm-pore-size membrane filter (polytetrafluoroethylene, DISMIC-25HP, Toyo Roshi Kaisha, Tokyo, Japan) to remove the 207 208 AC, and then the adsorbate concentration in the filtrate was measured. Complete removal of SSPAC required two filtrations. Preliminary tests confirmed that MIB did not adsorb on 209 the filter and that SSPAC was completely removed by the filtration. Solid-phase 210 211 concentrations of adsorbates were calculated from mass balances.

212

#### 213 **3. Results and discussion**

214

#### 215 3.1 SPAC and SSPAC production by normal micromilling

SSPACs with D50s of 137–198 nm were produced by the milling methods described in sec. 2.1, and SSPAC production with good reproducibility was verified both from the particle size distribution (Fig. S1) and by scanning electron microscopy (Fig. 2). The characteristics of the normally milled ACs are listed in Table S1.

220

221 3.2 Effects of particle size on MIB adsorption capacity

Equilibrium MIB adsorption isotherms were determined for Carbons A with D50s of 12.0–0.198  $\mu$ m (Fig. S2). The MIB adsorption capacity of each AC was quantified in terms of the solid-phase concentrations (designated as q) in equilibrium with liquid-phase concentrations of 50 and 100 ng/L, after fitting of the isotherm data to the Freundlich model. The effects of particle size on MIB adsorption capacity are depicted in Fig. 3a. Reduction of D50 from 12.0  $\mu$ m (PAC-A) to 1.47  $\mu$ m (SPACs-A-N) doubled the adsorption capacity, but the MIB adsorption capacity did not increase when D50 was decreased further: the MIB adsorption capacity of SSPAC-A-N (D50: 198 nm) was ~85% of that of SPACs-A-N.

A decrease in adsorption capacity in going from SPACs to SSPAC was also observed 230 231 for the other ACs. For example, the adsorption capacity of SSPAC-B-N was ~90% of that of SPACs-B-N, the adsorption capacity of SSPAC-C-N was 51-93% of that of SPACs-C-232 N, and the adsorption capacity of SSPAC-D-N was ~90% of that of SPACs-D-N (Fig. 3b-233 d, respectively). The adsorption capacity of SSPAC-D-N<sub>ET</sub>, which was produced by milling 234 for extended time, was 65-75% of that of SPACs-D-N. In the case of Carbons E, the 235 adsorption capacity started to decrease with particle size when D50 was  $<3.90 \mu m$ ; the 236 adsorption capacity of SSPAC-E-N was ~74% of that of SPAC<sub>L</sub>-E-N (Fig. 3e). A similar 237 238 trend was reported in a study of atrazine removal with microfiltration membrane precoated with SPAC (Amaral et al. 2016). In that study, adsorption capacity decreased as D50 was 239 reduced from ~300 to ~200 nm for one of the three studied ACs and as D50 was reduced 240 from  $\sim 600$  to  $\sim 500$  nm for another AC, although the authors of the paper did not comment 241 on the trend. 242

According to the SAM, adsorbent particle size affects MIB adsorption capacity when 243 the adsorbent radius is substantially larger than the adsorbate penetration depth (Matsui et 244 al. 2013, Matsui et al. 2014). Under these circumstances, the increase in the external surface 245 area of the adsorbent particles with decreasing particle size results in increased adsorption 246 247 capacity. Therefore, the higher adsorption capacity of PAC-A (12.0  $\mu$ m) relative to that of SPACs-A-N (1.47 µm) was likely due to this mechanism. That the adsorption capacity did 248 not increase with decreasing particle size for D50s of <1.47 µm (i.e., from SPACs-A-N to 249 250 SSPAC-A-N) suggests that the particle radii of both of these ACs were smaller than the

MIB penetration depth and that all of the internal pores of the particles were utilized for 251 MIB adsorption. Once the internal pores are completely utilized, adsorption capacity is 252 independent of particle size. The change in the AC particle size caused by micromilling 253 would not result in any substantial change in the internal pore structure (Ando et al. 2010, 254 Matsui et al. 2014, Pan et al. 2017). Moreover, any structural changes in internal pores 255 could not explain the phenomena that MIB adsorption capacity decreased while PSS-1100 256 257 adsorption capacity increased, as described in the section 3.6. Under these circumstances, pore surface chemistry strongly would affect adsorption capacity (Li et al. 2002, Matsui et 258 259 al. 2015, Quinlivan et al. 2005).

260

261 3.3 Oxygen content of AC

Oxygen-containing complexes on the AC surface alter surface hydrophobicity and affect the adsorption capacity for organic compounds, including MIB (Moreno-Castilla 2004, Pendleton et al. 1997, Song et al. 2010). The number of oxygen-containing complexes is reflected by the AC oxygen content (% [w/w]), which is thus highly correlated with MIB adsorption capacity (Li et al. 2002, Matsui et al. 2015).

The oxygen content of Carbons A increased with decreasing particle size (Fig. 4a), and 267 the increase was particularly large when D50 dropped to  $<1 \mu m$ . The external surface of 268 AC particles is usually more oxidized than the interior because oxygen diffuses very slowly 269 270 in the abundant, narrow internal pores (Boehm 2002). Micromilling in the bead mill, which was used to produce particles with D50s of  $<1 \mu m$ , brought the particle interior into contact 271 with oxygen in the water, possibly hastening oxidation. That the drop in MIB adsorption 272 capacity was coincident with a substantial increase in oxygen content (Figs. 3 and 4) 273 suggests that the drop in MIB adsorption capacity was due to carbon oxidation. 274

Wood-based Carbons A-C (i.e., PAC-A, PAC-B, and PAC-C) were produced by the 275 same manufacturer, and were marketed under the same specification and brand name. 276 However, they were produced in different batches, and were received in our laboratory at 277 different times. In addition, the times from receipt to experimental use (stock periods) 278 differed, decreasing in the order PAC-A (~7 years) > PAC-B (~1 year) > PAC-C (<6 279 months). This order is consistent with the order of decreasing oxygen content: PAC-A 280 (9.1%) > PAC-B (7.2%) > PAC-C (2.0%) (Fig. 4a–c). This finding is in line with a report 281 that AC is oxidized very slowly in air (Billinge et al. 1984, Boehm 2002). For these three 282 283 PACs, the order of decreasing MIB adsorption capacity was opposite the order of decreasing oxygen content: that is, adsorption capacity decreased with carbon age in the 284 order PAC-C > PAC-B > PAC-A (Fig. S2). Although characteristics other than oxygen 285 286 content might also be related to the adsorption capacity difference between the three carbons, these data is consistent with the idea that carbon oxidation decreased MIB 287 adsorption capacity (Considine et al. 2001). 288

For Carbons A-C, the sources of which had different oxygen contents, oxygen content 289 increased as the particle size was decreased by micromilling with the bead mill (Fig. 4a-c). 290 The same was true of Carbon D, a wood-based AC produced by a different manufacturer, 291 and Carbon E, a coal-based AC produced by a third manufacturer (Fig. 4d,e). The oxygen 292 content increased further when the milling time was extended from 1.5-2 to 5-7 h (compare 293 SSPAC-D-N and SSPAC-D-NET in Fig. 4d). Because the external surface of AC particles 294 295 is believed to be more oxidized than the interior (Boehm 2002, Partlan et al. 2016), overall particle oxidation would have increased as particle size was decreased by micromilling. 296 That oxygen content increased with decreasing particle size for all five kinds of ACs used 297 298 in this study is consistent with this understanding.

299

300 3.4 Controlling oxidation during SPAC and SSPAC production

The increase in oxygen content as milling progressed suggests that milling introduced oxygen-containing functional groups, which would increase the carbon hydrophilicity and thus decrease MIB adsorption capacity. Therefore, if oxidation during milling could be inhibited, the adsorption capacity decrease could be attenuated. We investigated various methods for producing less-oxidized SSPAC with D50s of <230 nm (Table S2, Fig. S1). The oxygen contents of Carbons C and D are presented in Fig. 5 (Fig. S3).

Oxidation-inhibiting milling (sec. 2.2.2) inhibited oxidation as D50 was reduced from 2.65-8.11  $\mu$ m to 174–230 nm. Specifically, oxidation-inhibiting milling to produce SSPAC-C-OI increased oxygen content by only 2.2 ± 0.5% (w/w) (Fig. 5c), whereas normal milling to produce SSPAC-C-N increased oxygen content by 3.4 ± 1.2% (Fig. 4c). The oxygen content increase in going from SPAC<sub>L</sub>-D to SSPAC-D by oxidation-inhibiting milling was 2.0 ± 0.2% (Fig. 5d), which was approximately half the increase due to normal milling (3.8 ± 0.1%; Fig. 4d).

We also measured the hydrogen, nitrogen, sulfur, and ash contents of ACs produced 314 by normal milling and oxidation-inhibiting milling (Fig. S4). Like oxygen content, 315 hydrogen content increased with decreasing particle size. When the particles were milled, 316 the ratio (increase in oxygen content, in moles)/(increase in hydrogen content, in moles) 317 ranged from 2/3 to 3/2. This result suggests that hydroxyl and/or carboxyl groups formed 318 319 on the carbon surface, an additional indication that milling increased carbon hydrophilicity. The ash contents in PAC-B-R, SPAC<sub>S</sub>-B-OI, SSPAC-B-OI, and SSPAC-B-N were 320 somewhat lower than those in PAC-B and SPACs-B-N. This result may have been due to 321 the effect of rinsing and dilution of the AC slurry with pure water; rinsing and dilution 322 would have washed out inorganic materials, including metals. However, this small change 323

in ash content did not affect MIB adsorption capacity: the MIB adsorption isotherm of PAC-

325 C-R did not differ substantially from that of PAC-C (Fig. S5).

326

327 3.5 Correlation between oxygen content and MIB adsorption capacity

We have suggested that the low MIB adsorption capacities of the SSPACs produced 328 by normal milling resulted from carbon oxidation. In a final test, SSPACs with low oxygen 329 content (SSPAC-C-OI) and with high oxygen content (SSPAC-C-N) were produced from 330 the same PAC, and their MIB adsorption capacities were determined. SSPAC-C-OI 331 332 exhibited higher adsorption capacity than SSPAC-C-N (Figs. S2 and S6). Moreover, there were strong negative correlations between oxygen content and MIB adsorption capacity 333 (Fig. 6). These results clearly indicate that the low MIB adsorption capacity of normally 334 335 milled SSPAC produced relative to that of SPACs was due to increased oxygen content, which increased carbon hydrophilicity. 336

The plots in Fig. 6 include data for AC particles with D50s from 137 nm to 1.64 µm 337 (i.e., all the SPACss and SSPACs). The correlation shown in the figure clearly means that 338 adsorption capacity was unrelated to particle size in this range, suggesting that the entire 339 340 particle, including internal pores, was accessible to MIB molecules, as suggested by the SAM (sec. 3.1). Oxygen content is a good indicator of MIB adsorption capacity of particles 341 with D50s of  $< \sim 1 \mu m$  (SPACs and SSPAC). This agrees with a previous finding that SPAC 342 hydrophilicity, as indicated by oxygen content, is the main determinant of adsorption 343 capacity for hydrophobic compounds such as geosmin (Matsui et al., 2015). 344

The oxygen content of SSPAC-OI was lower than that of SSPAC-N but was still higher than that of the SPACs from which SSPACs were produced. That is, oxidation was not completely prevented by oxidation-inhibiting milling. The highest adsorption capacity was attained with SPACs having a D50 of ~1  $\mu$ m. The energy required to reduce D50 to < ~1 µm would not offer any increase in adsorption capacity. Therefore, SPAC with a D50 of a few micrometers (~1.5  $\mu$ m for Carbons A–D, ~4.0  $\mu$ m for Carbon E) is a more reasonable choice for MIB removal from the viewpoint of high MIB adsorption capacity and reasonable energy consumption.

353

354 3.6 Effect of particle size on adsorption of other adsorbates

We observed that adsorption capacity for MIB, a hydrophobic low-MW compound 355 (log Kow, 3.31; 168 Da), increased in going from PAC to SPAC and decreased in going 356 357 from SPAC to SSPAC. To determine whether this trend held for other adsorbates, we conducted adsorption equilibrium tests on several environmentally relevant compounds 358 with various hydrophobicities and MWs. The adsorption capacities for these compounds 359 were quantified in terms of the solid-phase concentrations (q) in equilibrium with two 360 liquid-phase concentrations in the experimentally observed range (Fig. S6). Then we 361 evaluated the relationship between adsorption capacity and particle size (Fig. S8). 362

Going from SPACs-A-N (1.47 µm) to SSPAC-A-N (0.20 µm) did not decrease 363 adsorption capacity for PSS-1100 and SNOM, which have high MWs (>1000 Da, Fig. S8); 364 365 in fact, adsorption capacity consistently increased with decreasing particle size. This result is consistent with previous findings indicating that adsorption capacity for high-MW 366 compounds increases with decreasing AC particle size (Ando et al. 2010, Ando et al. 2011). 367 For high-MW compounds, the penetration depth was smaller than the radii of adsorbents 368 such as PAC and SPAC (Ando et al. 2010); therefore the interior of those adsorbents were 369 inaccessible to those adsorbates, and adsorption occurred only at the external particle 370 surface. In these circumstances, adsorption capacity increases with decreasing adsorbent 371 particle size. 372

For the other 7 compounds, all of which had MWs of <300 Da, adsorption capacity 373 decreased when D50 was decreased to  $< \sim 1 \mu m$ , as was the case for MIB. For all the 374 adsorbates, the adsorption capacities of SSPAC-A-N (0.20 µm) were lower than those of 375 SPACs-A-N (1.47 µm) (Figs. 7 and S8). The magnitude of the adsorption capacity decrease 376 depended on the adsorbate, but we were unable to identify the adsorbate property that 377 explained the magnitude of the decrease. For example, the magnitude of the decrease was 378 not correlated to adsorbate log Kow ( $R^2 < 0.02$ , Figs. 7 and S9a). The adsorption capacity 379 380 of SSPAC-A-N for geosmin (log Kow, 3.57), which is at least as hydrophobic as MIB (log Kow, 3.31), was 30% lower than that of SPACs-A-N. The adsorption capacity of SSPAC-381 A-N for DMTP-oxon (log Kow, 0.77), which is hydrophilic, was also 30% lower than that 382 of SPACs-A-N. For Carbons B (i.e., SPACs-B-N and SSPAC-B-N), the magnitude of the 383 adsorption capacity decrease was not correlated with adsorbate log Kow (Fig. S9b,c). The 384 lower adsorption capacity of SSPAC relative to that of SPACs was due to the higher 385 hydrophilicity of SSPAC, as indicated by the higher oxygen content (sec. 3.5). The 386 differences in the magnitude of the adsorption capacity decease in going from SPACs to 387 SSPAC for the eight compounds (MIB plus the seven adsorbates other than PSS-1100 and 388 SNOM) indicate that the adsorption capacities for these compounds had different 389 sensitivities to carbon hydrophilicity (oxygen content). 390

Different sensitivities to carbon hydrophilicity have been reported in other studies, although these studies did not involve SPAC or SSPAC. For example, a study comparing the adsorption capacities of AC for a hydrophobic compound, trichloroethene (log Kow, 2.42; 131 Da), and a hydrophilic compound, methyl *tert*-butyl ether (log Kow, 0.94; 88 Da), indicated that the decrease in adsorption capacity associated with increased carbon hydrophilicity was more pronounced for the hydrophilic compound (i.e., the adsorbate with the lower log Kow) (Li et al. 2002). However, a study involving several fatty acids,

including propanoic acid (log Kow, 0.33; 74.1 Da), valeric acid (log Kow, 1.39; 102 Da), 398 enanthic acid (log Kow, 2.42; 130 Da), and pelargonic acid (log Kow, 3.42; 158 Da), 399 revealed that the decrease in adsorption capacity associated with increased carbon 400 hydrophilicity was not necessarily greater for hydrophobic adsorbates with high log Kow 401 values than for hydrophilic adsorbates with low log Kow values (Kaneko et al. 1989). Water 402 likely adsorbs as clusters around the hydrophilic sites on the carbon surface (Pendleton et 403 al. 1997). As the number of hydrophilic sites increases, water clusters tend to extend over 404 hydrophobic sites, which prevents hydrophobic adsorbates, such as MIB, from accessing 405 406 the hydrophobic sites on the carbon surface. Increased formation of water clusters on hydrophilic sites also reduces the energy of interaction between hydrophilic adsorbates, 407 such as DMTP-oxon, and the carbon surface (Kaneko et al. 1989, Li et al. 2002). Although 408 409 adsorption capacity for both hydrophobic and hydrophilic adsorbates decreases with decreasing carbon hydrophobicity, the mechanisms of the decreases for the two types of 410 adsorbates are likely to differ, which in turn would lead to differences in the sensitivity of 411 adsorption capacity to carbon hydrophobicity. 412

413

#### 414 **4.** Conclusions

415 (1) Micromilling produced AC particles with D50s as low as 137 nm.

416 (2) MIB adsorption capacity decreased with decreasing particle size for ACs with D50s of
417 less than a few micrometers. This decrease in adsorption capacity was due to carbon
418 oxidization during micromilling.

419 (3) Partial inhibition of carbon oxidation during micromilling attenuated the decrease in
420 MIB adsorption capacity.

421	(4) For attaining high MIB adsorption capacity without unnecessary consumption of energy
422	for micromilling, SPAC with a D50 of 1–4 $\mu$ m is a reasonable choice.
423	(5) The adsorption capacity decrease with decreasing AC particle size for particles with
424	D50s of less than a few micrometers was observed for all eight low-MW adsorbates.
425	The magnitude of the decrease depended on the adsorbates, but the relationship could
426	not be explained simply by differences in adsorbate hydrophilicity.
427	(6) For SNOM and PSS-1100, adsorption capacity increased with decreasing adsorbent
428	particle size (i.e., in going from PAC to SSPAC).
429	(7) AC underwent oxidation during long-term storage, and the resulting increase in oxygen
430	content could be associated with decreases in adsorption capacity for compounds such
431	as MIB.
432	
433	Acknowledgments
434	
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441

440

research.

### 442 Appendix. Supplementary Information

Table S1-S3 and Figures S1–S9 are available in the online version at #######.

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- 589

 Table 1 – Commercial PACs used in this study.

					М	lanufacture	e specificatio	ns													
Designation	Series	Production year	Raw material	Raw Brand aterial name	Phenol number	Iodine number (mg/g)	Methylene blue number (mL/g)	ABS number	Manufacture												
PAC-A	Carbon-A	2008	wood	wood	wood	wood Taiko W						Futamura									
PAC-B	Carbon-B	2015					wood	Taiko W	< 25	> 900	> 150	< 50	Chemical Co.,								
PAC-C	Carbon-C	2016																			
PAC-D	Carbon-D	2016	wood	Shirasagi	< 25	> 900	> 150	< 50	Osaka Chemical Co., Osaka, Japan												
PAC-E	Carbon-E	2016	coal	6MD	< 25	> 900	> 150	< 50	Calgon Carbon Japan KK												



Fig. 1 – Schematic diagram of apparatus for bead milling under N<sub>2</sub>.



Fig. 2 – Scanning electron micrograph of SSPAC-B-N.





Fig. 3 – Plots of equilibrium MIB adsorption capacity versus AC particle size. Squares indicate PAC, diamonds indicate  $SPAC_L$ , triangles indicate  $SPAC_S$ , and circles indicate SSPAC. The adsorption capacities are given as solid-phase concentrations (q) at the equilibrium liquid-phase concentrations of 50 or 100 ng/L.



**Fig. 4** – **Plots of oxygen contents of Carbons A–E produced by normal milling versus AC particle size.** Squares indicate PAC, diamonds indicate SPAC<sub>L</sub>, triangles indicate SPAC<sub>S</sub>, and circles indicate SSPAC. Error bars, which indicate standard deviation, are obscured by the symbols.



**Fig. 5** – **Plots of oxygen contents of Carbons C and D produced by oxidation-inhibiting milling versus AC particle size.** Squares indicate PAC, diamonds indicate SPAC<sub>L</sub>, triangles indicate SPAC<sub>s</sub>, and circles indicate SSPAC. Error bars, which indicate standard deviation, are obscured by the symbols.



**Fig. 6** – **Correlation between MIB adsorption capacity and oxygen content.** Triangles indicate SPAC<sub>s</sub>, and circles indicate SSPAC.



Fig. 7 – Plot of ratios of SPAC<sub>s</sub> and SSPAC adsorption capacities versus adsorbate log Kow. The adsorption capacities were quantified in terms of the solid-phase adsorbate concentrations at equilibrium liquid-phase concentrations: MIB and geosmin, 50 ng/L; DMTP-oxon, 5  $\mu$ g/L; benzothiazole, phenol, *para*-nitrophenol (PNP), and salicylic acid, 0.5 mg/L; PSS-210, 2 mg/L. The ratio for PSS-210 is indicated by a dashed line because the log Kow is unavailable.

## Supplementary Information

# Effects of Decreasing Activated Carbon Particle Diameter from 30 μm to 140 nm on Equilibrium Adsorption Capacity

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Designation	Median diameter D50 (µm)	Mill	Source carbon	Median diameter D <sub>50</sub> (µm)	
SPAC <sub>L</sub> -A-N	not measured	Ball mill	PAC-A	12.0	
SPACs-A-N	1.47		SPAC <sub>L</sub> -A-N	not measured	
SSPAC-A-N	0.198	Bead mill	SPACs-A-N	1.47	
SPAC <sub>L</sub> -B-N	3.85	Ball mill	PAC-B	27.5	
SPACs-B-N	1.21		SPAC <sub>L</sub> -B-N	3.85	
SSPAC-B-N	0.194	Bead mill	SPACs-B-N	1.21	
SPAC <sub>L</sub> -C-N-1	not measured				
SPAC <sub>L</sub> -C-N-2	4.00				
SPAC <sub>L</sub> -C-N-3	3.87	Ball mill	PAC-C	23.1	
SPAC <sub>L</sub> -C-N-4	4.04				
SPAC <sub>L</sub> -C-N-5	3.79				
SPACs-C-N-1	1.12		SPAC <sub>L</sub> -C-N-1	not measured	
SPACs-C-N-2	0.949		SPAC <sub>L</sub> -C-N-2	4.00	
SPACs-C-N-3	1.64		SPAC <sub>L</sub> -C-N-3	3.87	
SPACs-C-N-4	1.30		SPAC <sub>L</sub> -C-N-4	4.04	
SPACs-C-N-5	not measured		SPAC <sub>L</sub> -C-N-5	3.79	
SPACs-C-N-6	1.45	Bead mill	SPAC <sub>L</sub> -C-N-6	4.44	
SSPAC-C-N-1	0.176		SPAC <sub>L</sub> -C-N-1	not measured	
SSPAC-C-N-2	0.137		SPAC <sub>L</sub> -C-N-2	4.00	
SSPAC-C-N-3	0.179		SPAC <sub>L</sub> -C-N-3	3.87	
SSPAC-C-N-4	0.177		SPAC <sub>L</sub> -C-N-4	4.04	
SSPAC-C-N-5	0.177		SPAC <sub>L</sub> -C-N-5	3.79	
SPAC <sub>L</sub> -D-N-1	3.03				
SPAC <sub>L</sub> -D-N-2	3.57	Ball mill	PAC-D	15.5	
SPAC <sub>L</sub> -D-N-3	3.00				
SPAC <sub>s</sub> -D-N-1	0.927		SPAC <sub>L</sub> -D-N-1	3.03	
SPACs-D-N-2	0.915		SPAC <sub>L</sub> -D-N-2	3.57	
SPACs-D-N-3	0.927		SPACs-D-N-3	3.00	
SSPAC-D-N-1	0.169	Dood mill	SPAC <sub>L</sub> -D-N-1	3.03	
SSPAC-D-N-2	0.164	Beau IIIII	SPAC <sub>L</sub> -D-N-2	3.57	
SSPAC-D-N-3	0.169		SPACs-D-N-3	3.00	
SSPAC-D-N <sub>ET</sub> -3	0.168		SPACs-D-N-3	3.00	
SSPAC-D-N <sub>ET</sub> -4	0.170		SPACs-D-N-3	3.00	
SPAC <sub>L</sub> -E-N	3.90	Ball mill	PAC-E	19.5	
SPAC <sub>S</sub> -E-N	1.19	Band mill	SPAC <sub>L</sub> -E-N	3.90	
SSPAC-E-N	0.177	Deau IIIII	SPAC <sub>L</sub> -E-N	3.90	

 Table S1 – Characteristics of normally milled PACs, SPACs, and SSPACs.

Table S2 – Characteristics of PACs, SPACs, and SSPACs produced by rinsing with pure water (R) and by oxidation-inhibiting milling (OI).

	Median		Detail	Production me			
Designation	diameter D <sub>50</sub> (µm)	Milling system	Rotational speed (rpm)	Bead/ball diameter (mm)	Slurry volume (mL)	Source carbon	
PAC-B-R	21.7		no			PAC-B	
SPAC <sub>L</sub> -B-OI	not measured	Ball mill – N <sub>2</sub> filled	45	5 & 10	100	PAC-B-R	
SPACs-B-OI	0.794	Bead mill –	2500	0.3	400 -	SPAC <sub>L</sub> -B-OI	
SSPAC-B-OI	0.230	N <sub>2</sub> purging	2390	0.1	500	SPAC <sub>L</sub> -B-OI	
PAC-C-R	22.7		no			PAC-C	
SPAC <sub>L</sub> -C-OI-1	not measured						
SPAC <sub>L</sub> -C-OI-2	5.15				100	PAC-C-R	
SPAC <sub>L</sub> -C-OI-3	3.82	Ball mill –	45	5 & 10			
SPAC <sub>L</sub> -C-OI-4	3.88	N <sub>2</sub> filled	45				
SPAC <sub>L</sub> -C-OI-5	8.11						
SPAC <sub>L</sub> -C-OI-6	2.78						
SPACs-C-OI-1	1.13			0.2		SPAC <sub>L</sub> -C-OI-1	
SPACs-C-OI-2	0.831				0.3		SPACL-C-OI-2
SSPAC-C-OI-3	0.213	Bead mill –	2500	0.1	400 -	SPAC <sub>L</sub> -C-OI-3	
SSPAC-C-OI-4	0.200	N <sub>2</sub> purging	2390	0.2	500	SPAC <sub>L</sub> -C-OI-4	
SSPAC-C-OI-5	0.229			0.3		SPAC <sub>L</sub> -C-OI-5	
SSPAC-C-OI-6	0.174			0.1	1	SPAC <sub>L</sub> -C-OI-6	
PAC-D-R	15.3	no				PAC-D	
SPAC <sub>L</sub> -D-OI-1	3.87						
SPAC <sub>L</sub> -D-OI-2	2.65	Ball mill – N2 filled	45	5 & 10	100	PAC-D-R	
SPAC <sub>L</sub> -D-OI-3	3.21						
SSPAC-D-OI-2	0.197	Bead mill –	2500	0.1	400 -	SPAC <sub>L</sub> -D-OI-2	
SSPAC-D-OI-3	0.197	N <sub>2</sub> purging	2390	0.1	500	SPAC <sub>L</sub> -D-OI-3	

 Table S3 – Composition of the working solution.

Ion	Na <sup>+</sup>	$\mathrm{K}^{\scriptscriptstyle +}$	Mg <sup>2+</sup>	Ca <sup>2+</sup>	Cl-	NO <sup>3-</sup>	SO4 <sup>2-</sup>	Alkalinity
Concentration (mg/L)	19	3.1	24	18	31	6.9	14	50





Fig. S1 – Particle size distributions of PACs, SPACs, and SSPACs.





**Fig. S2 – MIB adsorption isotherms for Carbons A–E.** Squares indicate PAC, diamonds indicate SPAC<sub>L</sub>, triangles indicate SPAC<sub>S</sub>, and circles indicate SSPAC.



Fig. S3 –Plots of oxygen contents of Carbons C and D produced by normal and oxidation-inhibiting millings versus AC particle size. Squares indicate PAC, diamonds indicate SPACL, triangles indicate SPACS, and circles indicate SSPAC. Closed plots indicate carbons produced by normal milling and open plots indicate those by oxidation-inhibiting milling. Error bars, which indicate standard deviation, are obscured by the symbols.



Fig. S4 – Hydrogen, nitrogen, sulfur, oxygen, and ash contents of Carbons-B produced by normal milling and oxygen-inhibiting milling.



Fig. S5 – MIB adsorption isotherms of PAC-C-N and PAC-C-R (produced by rinsing PAC-C-N with pure water).



**Fig. S6** – **MIB adsorption isotherms for SPAC**<sub>L</sub>, **SPAC**<sub>S</sub>, and **SSPAC prepared from Carbons C and D by oxidation-inhibiting milling.** Diamonds indicate SPAC<sub>L</sub>, triangles indicate SPAC<sub>S</sub>, and circles indicate SSPAC.





**Fig. S7** – **Isotherms for adsorption of 9 adsorbates on Carbons A.** Squares indicate PAC, triangles indicate SPACs, and circles indicate SSPAC.





**Fig. S8** – **Plots of equilibrium adsorption capacity versus particle size (Carbon A) for 9 adsorbates.** Squares indicate PAC, triangles indicate SPAC<sub>s</sub>, and circles indicate SSPAC.



Fig. S9 – Plots of ratios of SPACs and SSPAC adsorption capacities versus adsorbate log Kow. Adsorbate concentrations I and II (panels b-1 and b-2, respectively): MIB and geosmin, 50 and 100 ng/L; DMTP-oxon, 5 and 10  $\mu$ g/L; benzothiazole, phenol, *para*-nitrophenol (PNP), and salicylic acid, 0.5 and 1 mg/L; PSS-210, 2 and 4 mg/L. The ratios for PSS-210 are indicated by the dashed lines because the log Kow value is unavailable.