

Supporting Information

An Active Absorbent for Cleanup of High-Concentration Strong Acid and Base Solutions

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Experimental section

Preparation of MFS/Fe₃O₄

FeCl₃·6H₂O (0.9 g) and CH₃COONa (2.4 g) were dissolved in ethylene glycol (30 mL) under vigorous stirring for 5 min at room temperature. The resulting solution was transferred to a Teflon-lined autoclave bomb and heated in an oven at 200 °C for 8 h. The final product was washed 3 times with ethanol and DI water. A mixture containing EtOH (100 mL) and MNPs (200 mg) was stirred for 1 min. Then, an MFS (2 × 2 × 2 cm³) was added to the mixture, followed by ultrasonication for 10 min. After 10 min, the resulting sponge was dried in an oven at 50 °C without washing. The resulting MFS/MNP was washed 3 times with DI water and dried in an oven at 50 °C.

Preparation of MFS/Au or Ag

Gold(III) chloride trihydrate or silver(I) nitrate (0.066 g) was dissolved in EtOH (17 mL) under vigorous stirring for 5 min at room temperature. Then, an MFS (2 × 2 × 2 cm³) was added to the resulting solution, followed by stirring for 4 h. After 4 h, the resulting sponge was washed 3 times with DI water. The resulting sponge was dipped in NaBH₄ (10 mM) solution for 30 min, washed 3 times with DI water, and dried in an oven at 50 °C.

Preparation of MFS/TiO₂

Titanium tertbutoxide (7.1 mL) was dissolved in EtOH (50 mL) under vigorous stirring for 1 min at room temperature. Then, 1 mL of DI water was added to the solution, which was stirred for 30 min, and 0.5 mL of HCl solution (pH=3) was added to the resulting solution. An MFS (2 × 2 × 2 cm³) was added to the resulting solution, followed by stirring for 4 h. After 4 h, the resulting sponge was washed 3 times with DI water and dried in an oven at 50 °C.

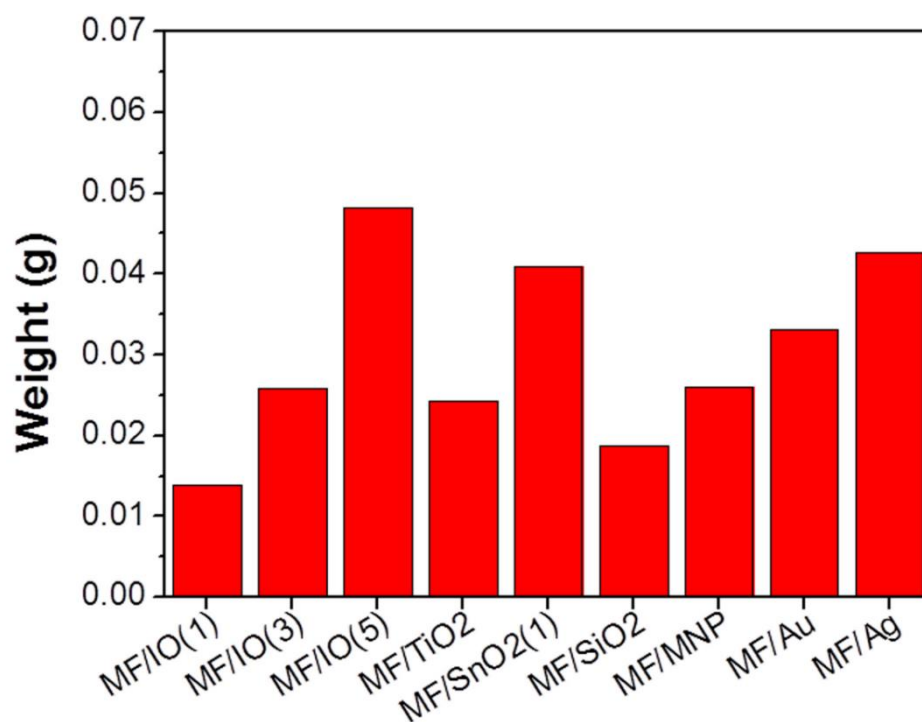


Figure S1. Weight data of various absorbents. All absorbents had a same dimensional size and volume. The measurements were repeated three times and average values were used.

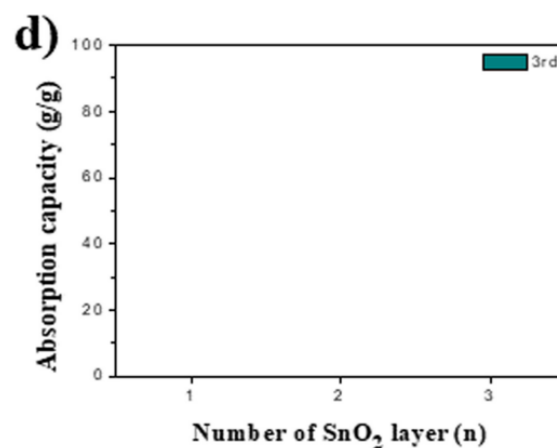
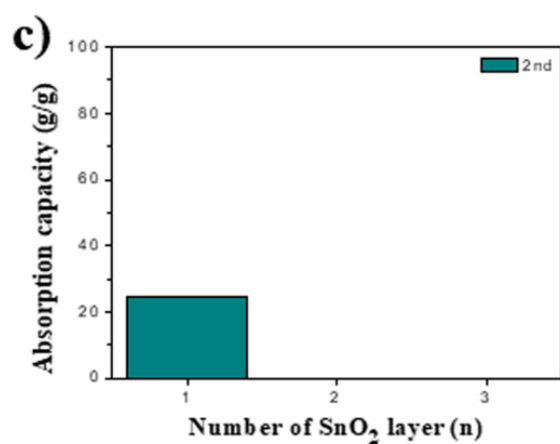
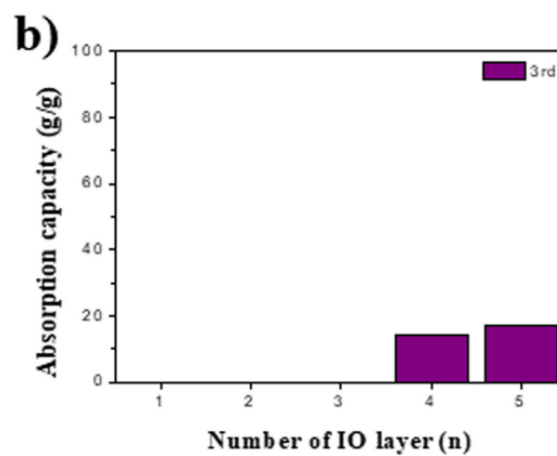
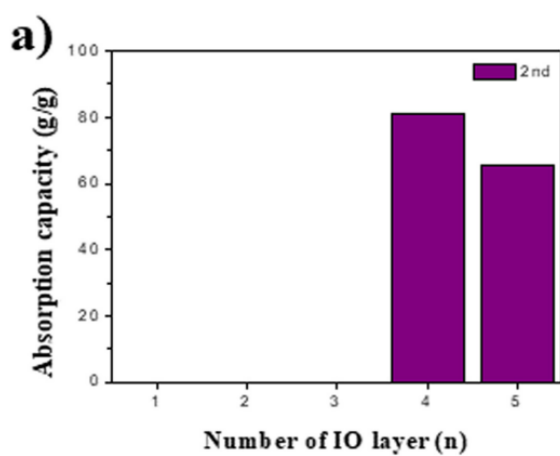


Figure S2. Recyclability test of (a,b) MFS/IO(1–5) and (c,d) MFS/SnO₂(1–3) for (a,c) 2nd and (b,d) 3rd use for the absorption of a NaOH solution (50%).

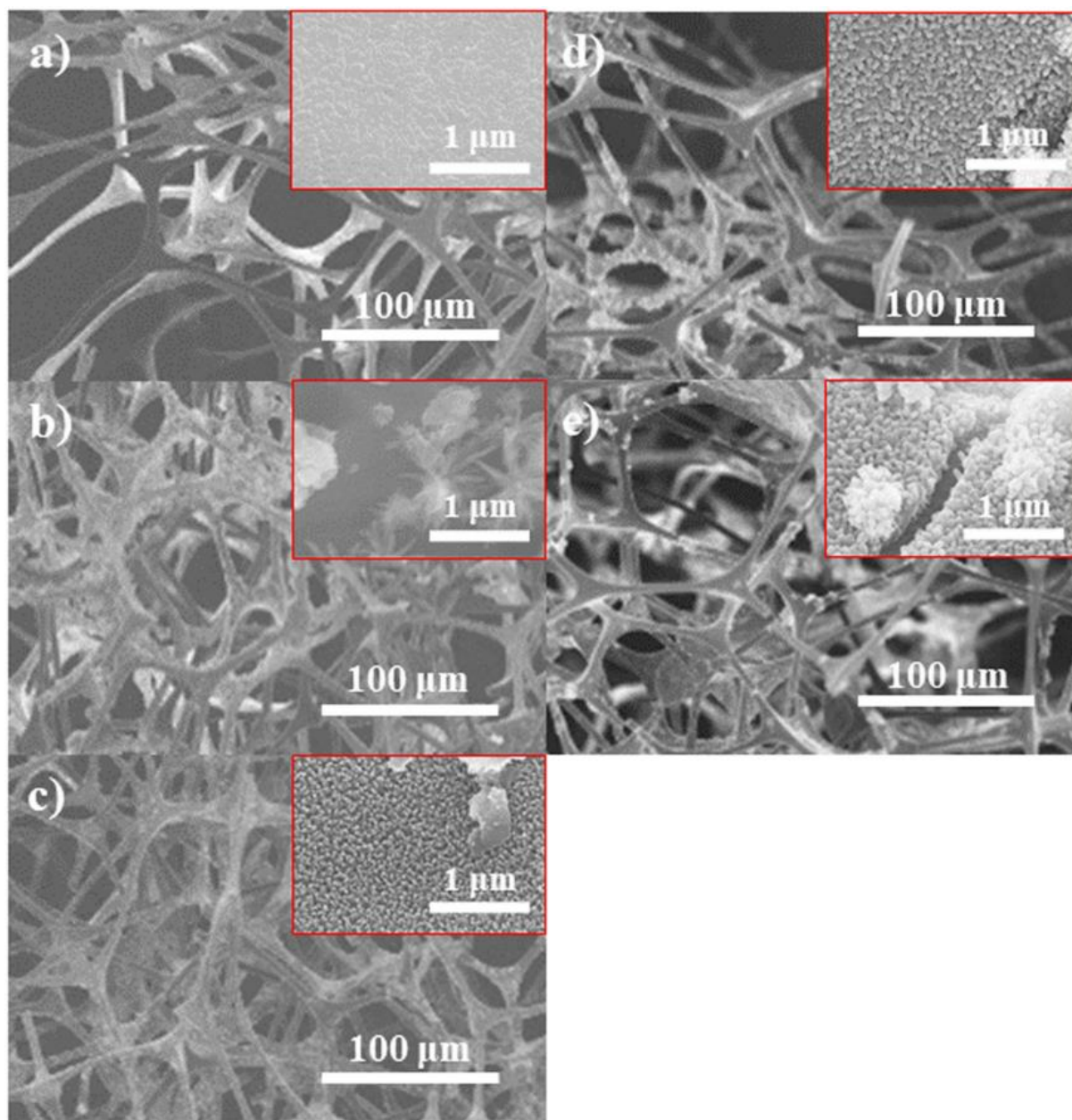


Figure S3. SEM images of MFS/IO(1–5) after the 1st absorption of a NaOH solution (50%). (a) MFS/IO(1), (b) MFS/IO(2), (c) MFS/IO(3), (d) MFS/IO(4), and (e) MFS/IO(5).

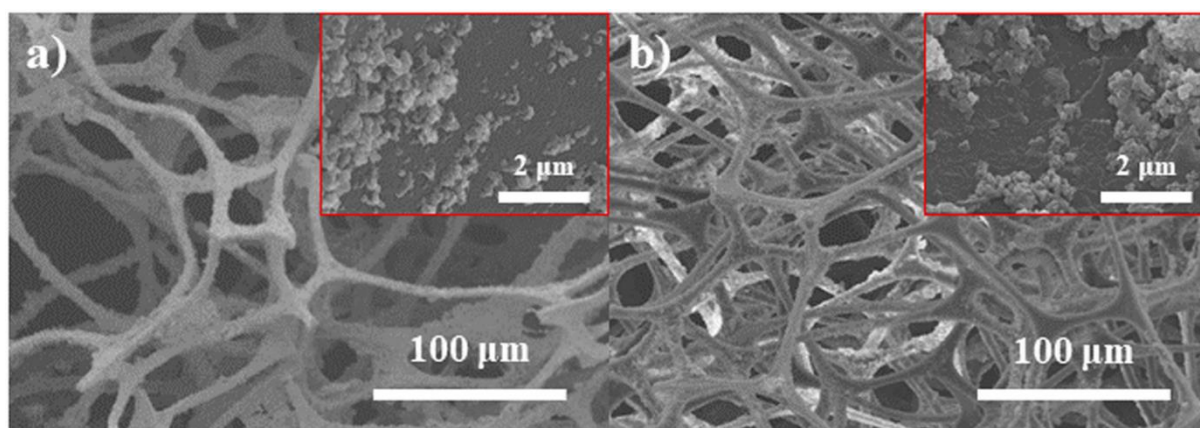


Figure S4. SEM images of MFS/Fe₃O₄ (a) before and (b) after dropping MFS/Fe₃O₄ on a NaOH solution (50%).

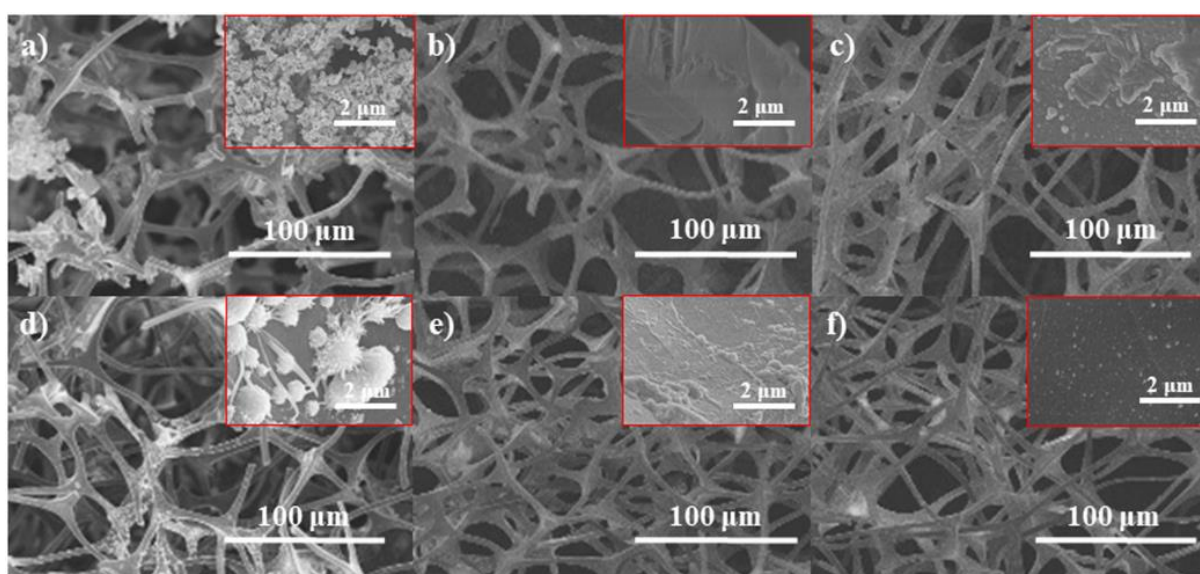


Figure S5. SEM images of MFS/SnO₂ (a–c) before and (d–f) after the 1st absorption cycle of a NaOH solution (50%). (a,d) MFS/SnO₂(1), (b,e) MFS/SnO₂(2), and (c,f) MFS/SnO₂(3).

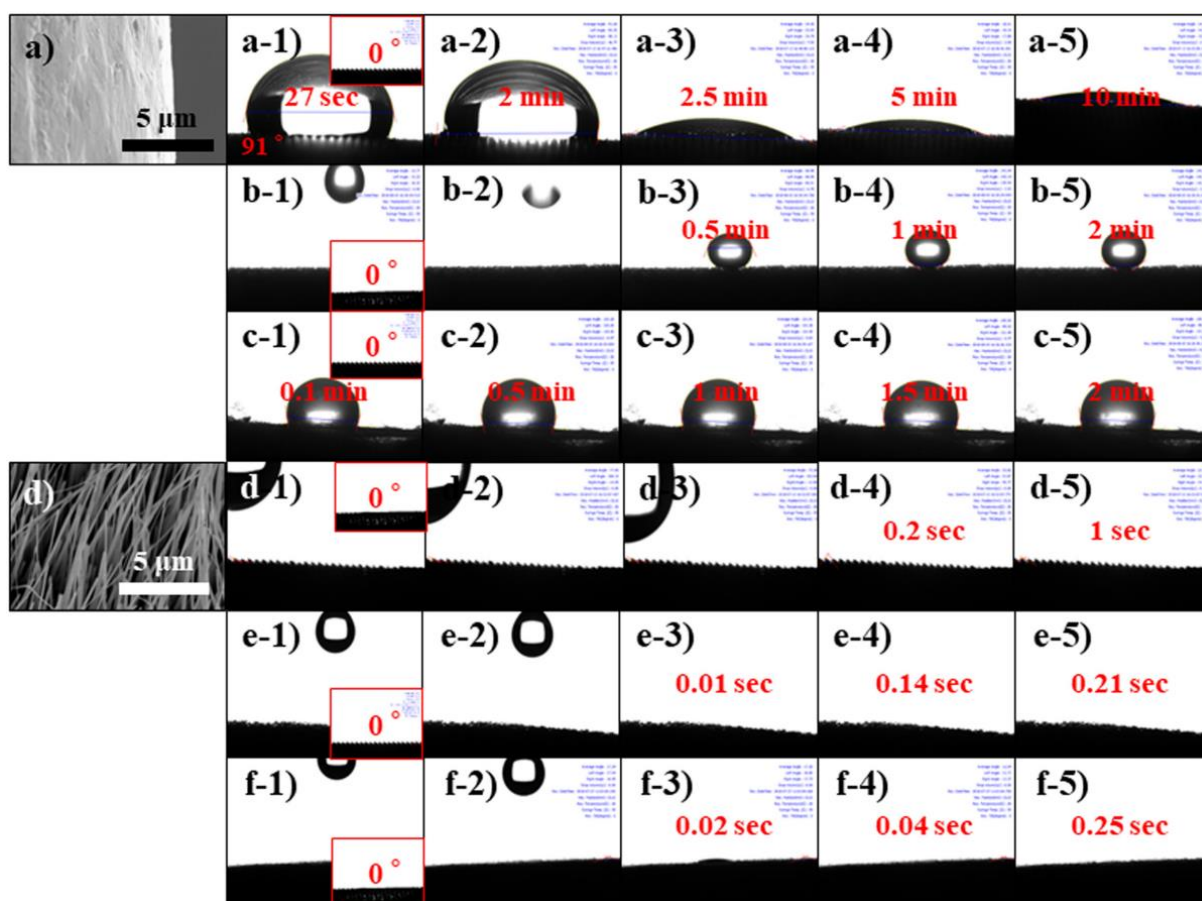


Figure S6. Contact angle and absorption time data of a droplet of 50% NaOH solution upon dropping a droplet on various absorbents or substrates. (a1–a5) Superhydrophilic Cu mesh with a smooth surface and (a) its SEM image, (b1–b5) MFS, (c1–c5) absorption pad, (d1–d5) superhydrophilic Cu mesh with a needlelike surface and (d) its SEM image, (e1–e5) MFS/IO(1), and (f1–f5) MFS/IO(5). Each inset shows the WCA of each absorbent or substrate.

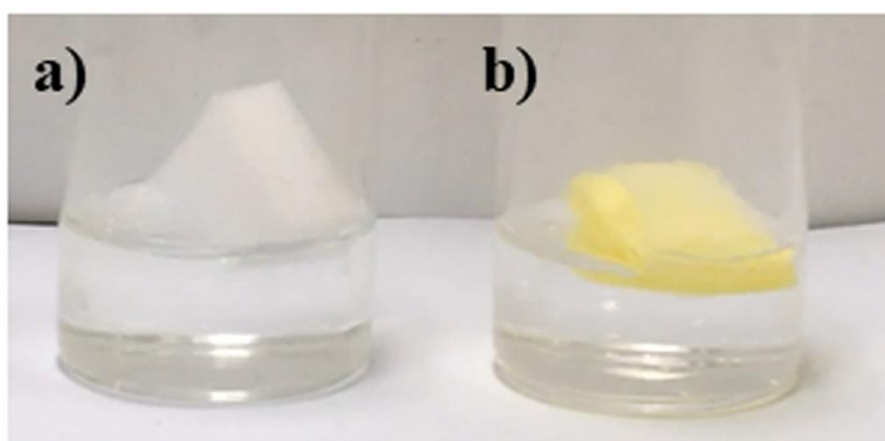


Figure S7. Drop tests of absorbents on a HCl solution (15%). (a) MFS and (b) absorption pad.

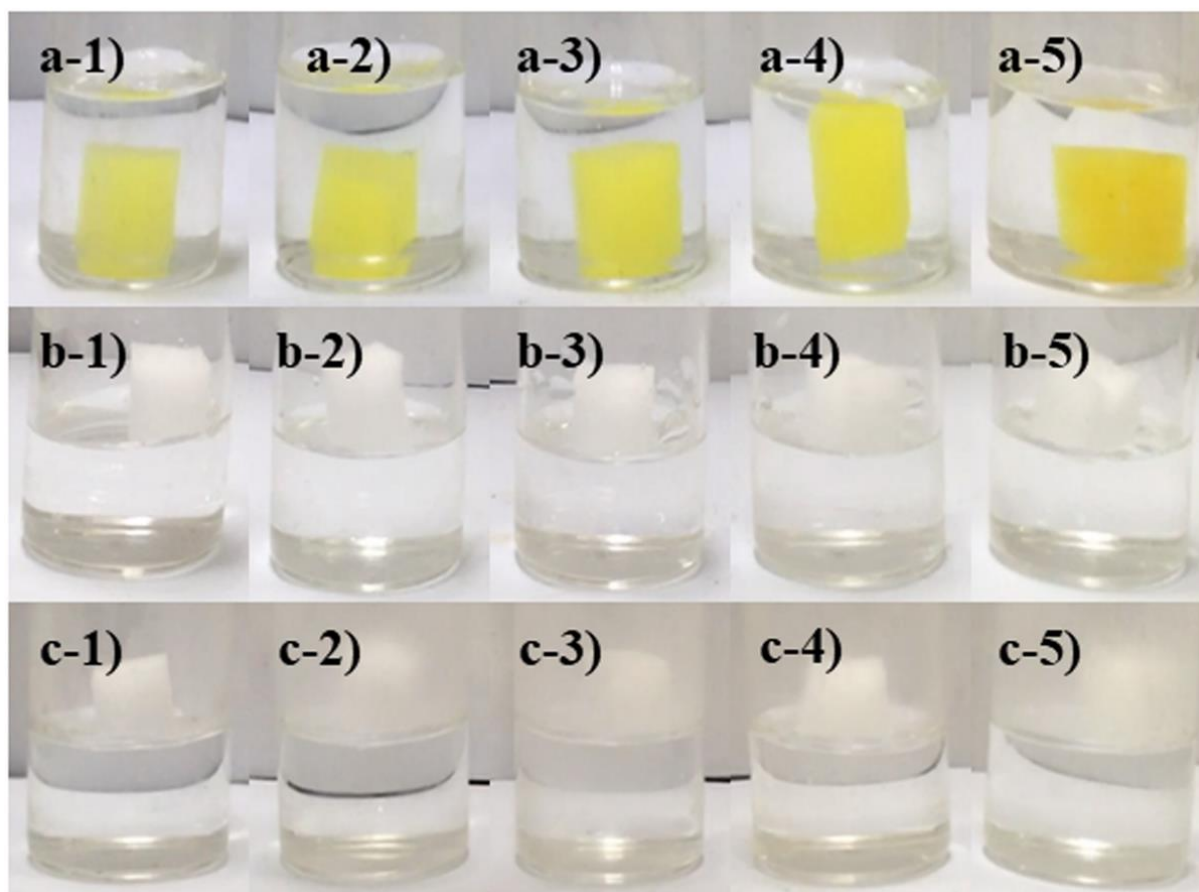


Figure S8. Recyclability test of MFS/IO(1–5) for absorption of a HCl solution (15%). (a1–a5) 1st absorption, (b1–b5) 2nd absorption, and (c1–c5) 3rd absorption cycle. (a1,b1,c1) MFS/IO(1), (a2,b2,c2) MFS/IO(2), (a3,b3,c3) MFS/IO(3), (a4,b4,c4) MFS/IO(4), and (a5,b5,c5) MFS/IO(5).