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# **Supporting Information**

# One-dimensional controllable crosslinked polymers grafted with N-methyl-D-

## glucamine for effective boron adsorption

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Fig. S1 TEM images of polymers with different mass ratios



**Fig. S2** N<sub>2</sub> adsorption/desorption isotherms and pore size distribution (a, b) Poly(2/1) and Poly(2/1)-NMDG, respectively; and (c, d) Poly(5/1) and Poly(5/1)-NMDG), respectively.



**Fig. S3** SEM images of Poly(5/1)-NMDG before sorption (a) and after four cycles (b); Poly(6/1)-NMDG before sorption (c) and after four cycles (d)

#### Part. S1

During the regeneration of adsorbents, Poly(5/1)-NMDG (0.7 g) and Poly(6/1)-NMDG (0.7 g) were added into 70 mL of 500 mg L<sup>-1</sup> H<sub>3</sub>BO<sub>3</sub> solution at pH=8, this solution was oscillated (200 rpm) for 24 h at room temperature. After filtration, the solution and sorbents were collected, respectively. The loaded sorbents were eluted with 70 mL HCl (1 mol L<sup>-1</sup>) and NaOH (1 mol L<sup>-1</sup>) solution, respectively. After filtration, the adsorbents were washed to neutrality with water and alcohol, and dried in oven for 24 h. Then, the regenerated sorbents were reused in the next adsorption experiments.

To study the influence of pH on boron removal, 250 mg  $L^{-1}$  H<sub>3</sub>BO<sub>3</sub> solutions with different pH values (4, 5, 6, 7, 8, 9, 10±0.05) were prepared. The Poly(5/1)-NMDG adsorbent (0.1 g) was added into the 10 mL H<sub>3</sub>BO<sub>3</sub> solution with different pH values, respectively. The mixture was oscillated (200 rpm) for 24 h at room temperature.

For adsorption isotherm experiments, the sorbent (0.1 g) was added into 10 mL different concentration boric acid (5, 10, 20, 50, 100, 200, 250, 350, 500 mg L<sup>-1</sup>) solutions at pH=9, room temperature. The mixture reacted for 24 h. The boron concentration of the filtrate was measured once the equilibrium reached.

The adsorption kinetic experiments were carried out by adding 0.1g sorbents into the 10 mL boric acid solution (350 mg L<sup>-1</sup>) at pH=9. They were oscillated (200 rpm) for different time (10, 20, 30, 60, 120, 240, 480, 720 min) at room temperature. After the filtration, all the liquids were tested to obtain the boron concentration.

## Part. S2

Langmuir eqn (2) and Freundlich eqn (3) are shown below:

$$\frac{C_e}{Q_e} = \frac{1}{Q_m}C_e + \frac{1}{K_L Q_m}$$
(2)

$$lgQ_e = \frac{l}{n} lgC_e + lgK_F$$
(3)

Here,  $C_e$  is the equilibrium concentration,  $Q_e$  denotes the equilibrium adsorption capacity,  $Q_m$  is the maximum adsorption capacity,  $K_L$  represents the Langmuir isothermal constant,  $K_F$  and n are Freundlich constants related to adsorption capacity and adsorption rate.



Fig. S4 Isotherm adsorption curves of the Langmuir (a) and Freundlich (b) models

## Part. S3

Pseudo-first-order kinetic eqn (4) and pseudo-second-order kinetic eqn (5):

$$lg(Q_{e} - Q_{t}) = lgQ_{e} - \frac{k_{1}}{2.303}t$$

$$\frac{t}{Q_{t}} = \frac{1}{k_{2}Q_{e}^{2}} + \frac{1}{Q_{e}}t$$
(4)
(5)

Here,  $Q_e$  is equilibrium capacity,  $Q_t$  is boron adsorption at different time,  $k_1$  and  $k_2$  are pseudofirst-order kinetics and pseudo-second-order kinetics adsorption rate constant respectively.



Fig. S5 Fitting curves of the pseudo-first-order (a) and pseudo-second-order (b) kinetics

Table S1. Adsorption conditions and adsorption capacity of magnetic boron adsorbents

Sample	PH	C <sub>0</sub> (mg L <sup>-1</sup> )	t (min)	T (°C)	Adsorption (mg g <sup>-1</sup> )
Poly(2/1)@Fe <sub>3</sub> O <sub>4</sub> -NMDG	8	350	120	20	5.4