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Supporting Information

Facile synthesis of thiazole-functionalized magnetic microspheres for

highly specific separation of heme proteins

Binghai Wang^a, Juanqiang Wang^a, Qian Shao^a, Xingjun Xi^b, Qiao Chu^b, Genlai Dong^b and Yun Wei^a*



Fig. S1 HRTEM image of Fe₃O₄ microspheres

^{*} Corresponding author: Yun WEI, State Key Laboratory of Chemical Resource Engineering, Beijing University of Chemical Technology, 15 Beisanhuan East Road, Chaoyang District, Beijing 100029, China, Tel & fax: 0086 10 64442928. E-mail: <u>weiyun@mail.buct.edu.cn</u>



Fig. S2 XRD patterns of Fe₃O₄ (a), Fe₃O₄@SiO₂ (b) and Fe₃O₄@SiO₂@AT (c)

The diffraction peaks (2θ =30.1°, 35.5°, 43.1°, 53.4°, 57.0° and 62.6°) for Fe₃O₄, Fe₃O₄@SiO₂, Fe₃O₄@SiO₂@AT were indexed as (220), (311), (400), (422), (511) and (440), respectively. The positions of these peaks matched well with database for magnetite in the JCPDS-International Center for Diffraction Data (JCPDS Card: 19-629) file. The XRD patterns indicated that the Fe₃O₄ crystalline structure did not change before and after each step of the chemical modification reaction.



Fig. S3 The size-distribution analysis of Fe₃O₄@SiO₂@AT with a log-normal fit



Fig. S4 Raman spectra of Fe₃O₄ microspheres



Fig. S5 Zeta potentials of Fe₃O₄@SiO₂@AT in different pH solutions



Fig. S6 The use of the recycled Fe₃O₄@SiO₂@AT for hemoglobin adsorption

The result of recycle experiment was shown in Fig. S6. The hemoglobin adsorption-desorption process was repeatedly used for 6 cycles, and the adsorption efficiency was still maintained at 85%. It indicated that the prepared microspheres were very stable for separation of hemoglobin.



Fig. S7 N₂ adsorption/desorption isotherms (at 77K) and the pore size distribution curves (inset) of the Fe₃O₄@SiO₂@AT and Fe₃O₄@SiO₂@IL composite

Table S1	Properties	of differen	t adsorbents	for	hemoglobin	capture

Adsorption material	Size	Capacity for	Capture time
		hemoglobin	
Mesoporous silica ⁶²	50 and 10 μ m (pore diameters	300 mg/g	30 min
	from 6 to 20 nm)		
Mesoporous TiO ₂ -SiO ₂ ⁶³	Wall thickness ~5 nm (pore	301.8 mg/g	12 h
	diameters from 5.8 to 7.25 nm)		
Fe ₃ O ₄ @SiO ₂ @IL(previous	~300 nm (pore diameter 13.71 nm)	~2.0 g/g	15 min
work) ⁴⁹			
Fe ₃ O ₄ @SiO ₂ @AT (This work)	322 nm (pore diameter 11.6 nm)	2.02 g/g	15 min

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