Supporting Information

Enantioselective Absorption of Enantiomers with Maleic Anhydride-β-

Cyclodextrin Modified Magnetic Microspheres

Jun Huang, Ping Su, Jingwei Wu, Yi Yang*

College of Science & Beijing Key Laboratory of Environmentally Harmful Chemical

Analysis, Beijing University of Chemical Technology, Beijing 100029, P. R. China.

*Author for correspondence: e-mail: yangyi@mail.buct.edu.cn

Fax: +86-10-64434898 Phone: +86-10-64454599

Synthesis of the β-CD monomer (MAH-β-CD)

The MAH- β -CD monomer, which can be copolymerized with other compounds containing a double bond, was completed by nucleophilic attack on the alcohol group in a dimethylformamide (DMF) solution. Typically, 0.005 mol of β -CD and 0.05 mol of MAH were dissolved in 30 mL of dried DMF under adequate ultrasonication, and the transparent mixture solution was subsequently placed in an 80°C oil bath for 10 h under vigorous stirring. After the reaction was completed and the mixture cooled, a yellow mixture solution was obtained. At this point, a white precipitate was immediately filtrated with the addition of 30 mL of trichloromethane, and washed with 100 mL of acetone four times. The final product was dried in a vacuum oven until a constant weight was maintained.

The elemental analytical result of the MAH- β -CD monomer is C: 45.68%; H: 5.19%, which indicate that the molecular structure of MAH- β -CD is agreement with the theoretic calculation of C₆₂H₈₀O₅₀. The degree of substitution is five. ¹³C-NMR (400MHz, D₂O as the solvent, Fig. S-1) δ (ppm): 169 (C-10), 166 (C-7), 132 (C-8), 127 (C-9).

References

Y.Y. Liu, X.D. Fan. Polymer 2002, 43,4997-5003.

Fig. S-1: ¹³C-NMR spectrum of MAH-β-CD monomer

