

*Supporting Information*

# Preparation of a Molecularly Imprinted Film on Quartz Crystal Microbalance Chip for Determination of Furanic Compounds

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## Supporting Information

### 3. Results and Discussion

#### 3.1. Synthesis and properties of (*Methacr-L-Cys-NHBn*)<sub>2</sub> and (*Methacr-L-Ser-NHBn*) Characterization of (*Methacr-L-Cys-NHBn*)<sub>2</sub>

<sup>1</sup>H NMR (300 MHz, CDCl<sub>3</sub>, δ): 1.77 (s, 3H), 3.02 (m, 2H), 4.4 (q, 2H), 5.22 (s, 1H), 5.5 (d, 2H), 7.22-7.28 (m, 5H).

<sup>13</sup>C NMR (75.5 MHz, CDCl<sub>3</sub>, δ): 18.3, 43.8-44, 46.7, 53.6, 121, 127.4-127.6, 128-128.2, 128.7, 137.6-137.8, 138.9, 168.5, 170.2

IR(cm<sup>-1</sup>):164.78, 2926.59, 3317.1

[α]<sub>D</sub><sup>28</sup> = +72° (CH<sub>2</sub>Cl<sub>2</sub>, c=0.325, l=1 dm)

Mass (MALDI) = 555.28 m/z

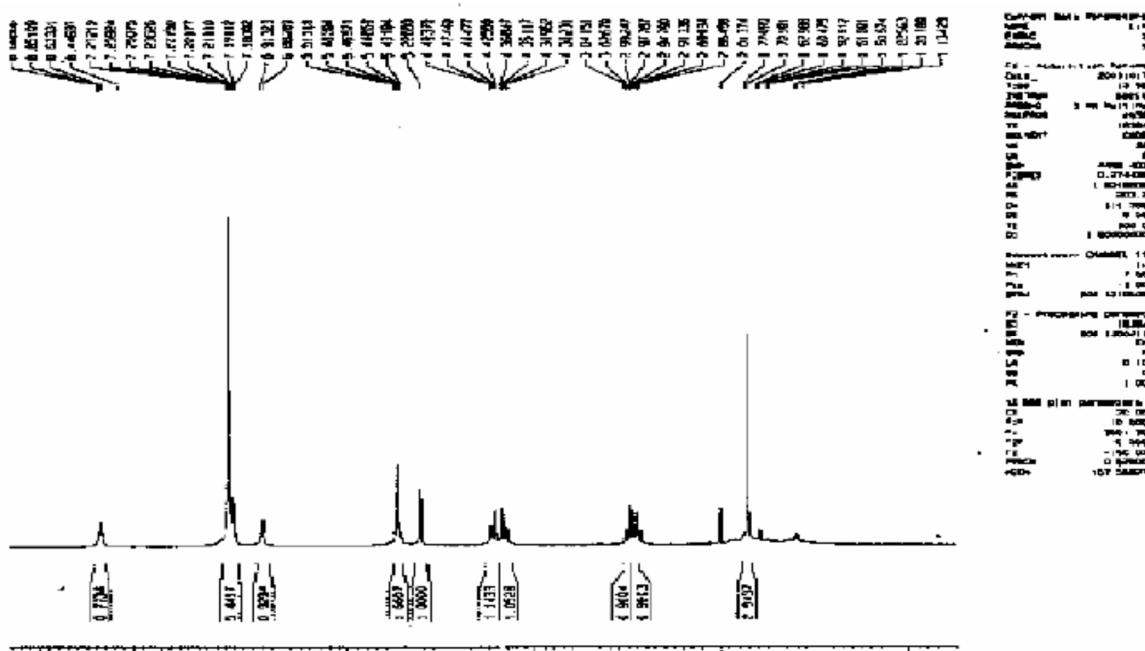
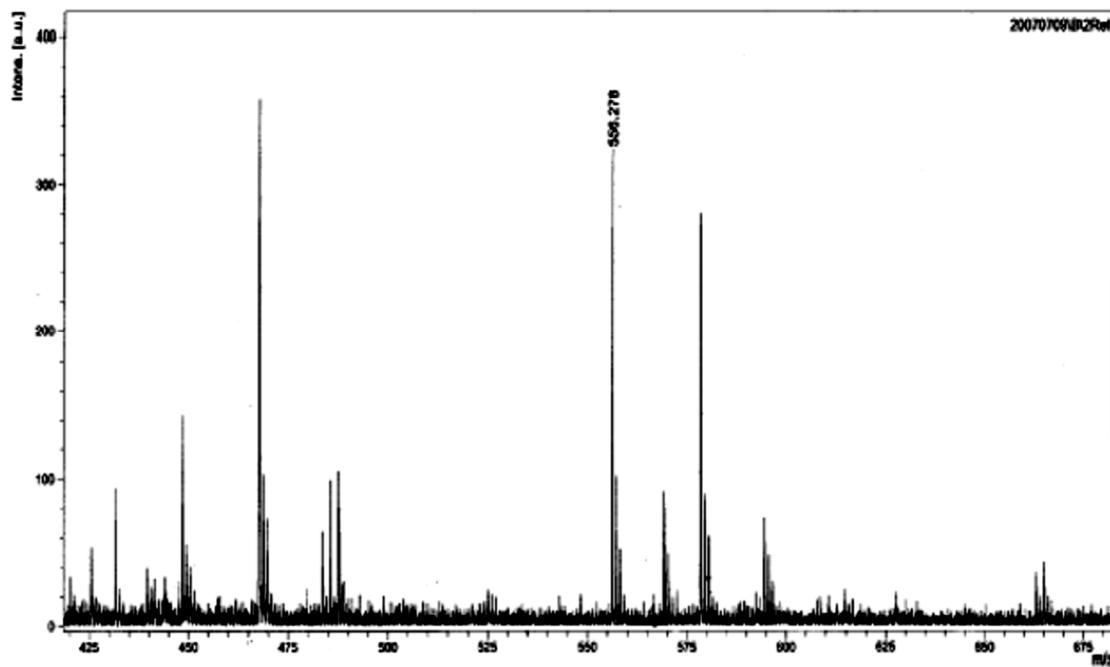
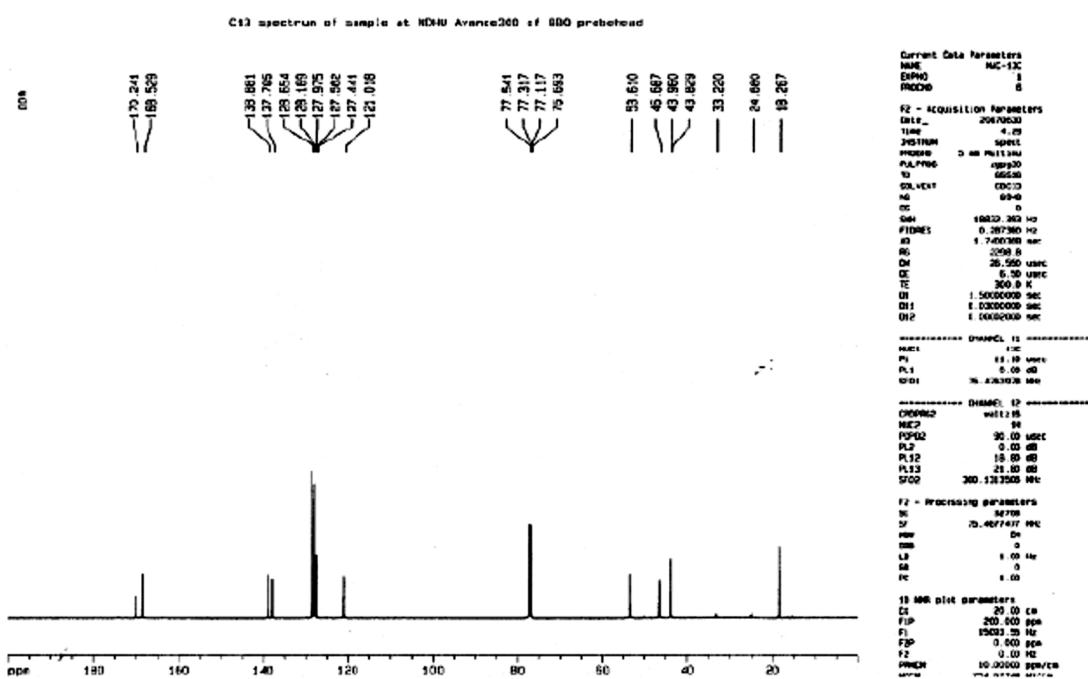
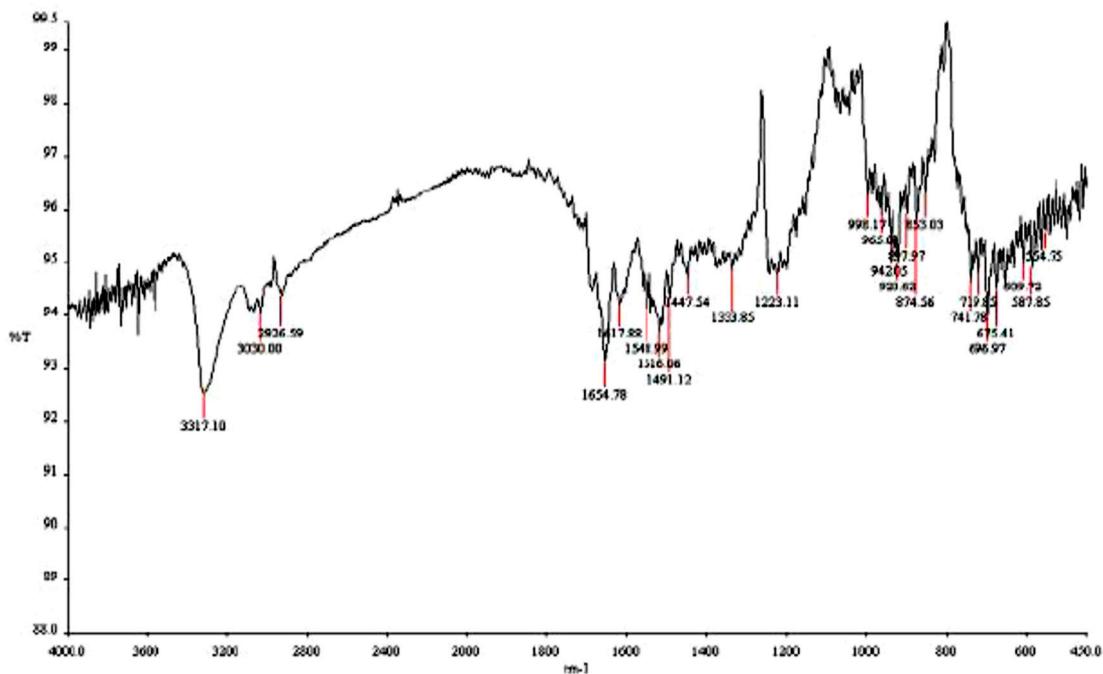


Figure S1. <sup>1</sup>H NMR spectrum of (Methacr-L-Cys-NHBn)<sub>2</sub>.



**Figure S2.**  $^{13}\text{C}$  NMR spectrum of (Methacr-L-Cys-NHBn)<sub>2</sub>.





**Figure S4.** IR spectrum of (Methacry-L-Cys-NHBn)<sub>2</sub>.

### 3.2. Characterization of Methacry-L-Ser-NHBn

<sup>1</sup>H NMR (300 MHz, MeOH-d<sub>4</sub>, δ): 1.95 (s, 3H), 3.82 (m, 2H), 4.44-4.46 (d, 2H), 4.56-4.59 (m, 1H), 5.43 (s, 1H), 5.81 (s, 1H), 7.24-7.34 (m, 5H).

<sup>13</sup>C NMR (75.5 MHz, MeOH-d<sub>4</sub>, δ): 17.3, 42.8-43.2, 55.6-55.8, 61.5-61.7, 120, 126.78-126.84, 127.01-127.1, 128.13, 138.3-139.5, 142.6, 169.68, 170.85-171.1

IR(cm<sup>-1</sup>): 1041.80, 1658.52, 2879.10-2939.32, 3313.70

$[\alpha]_D^{25} = -8.33^\circ$  (MeOH, c=0.3, l=1 dm)

Mass (MALDI) = 262.42 m/z

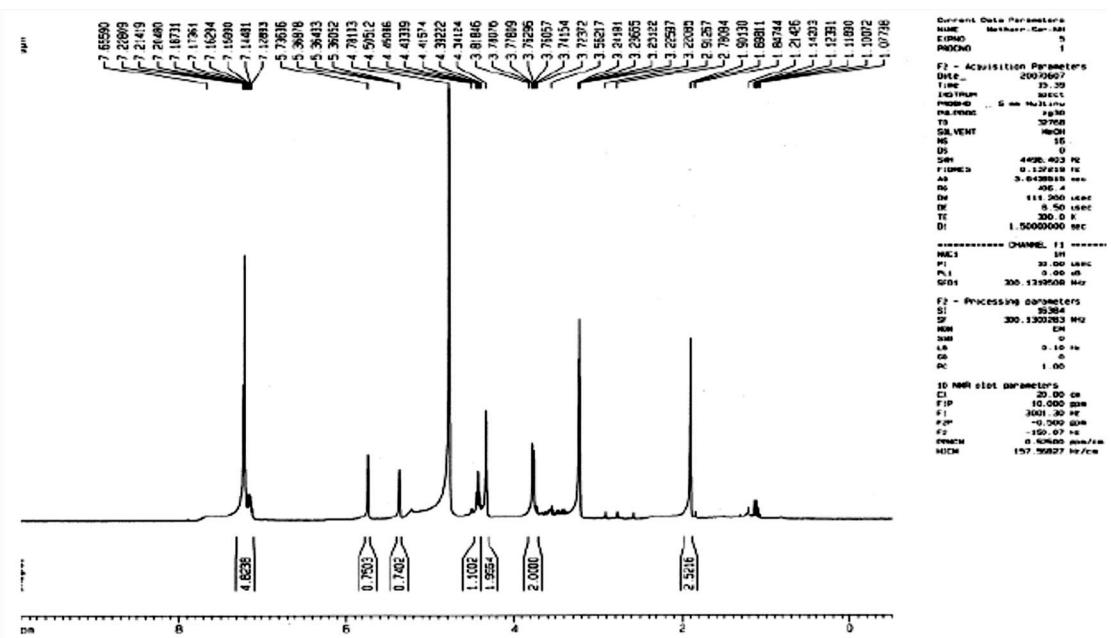


Figure S5.  $^1\text{H}$  NMR spectrum of Methacr-L-Ser-NHBn.

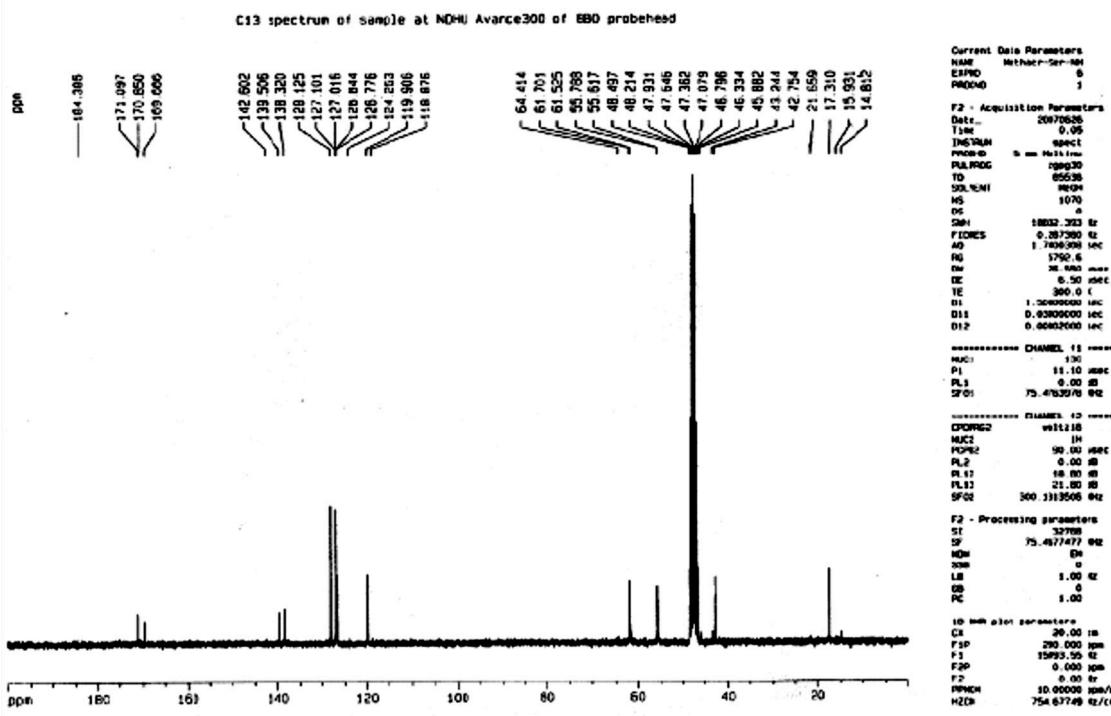


Figure S6.  $^{13}\text{C}$  NMR spectrum of Methacr-L-Ser-NHBn.

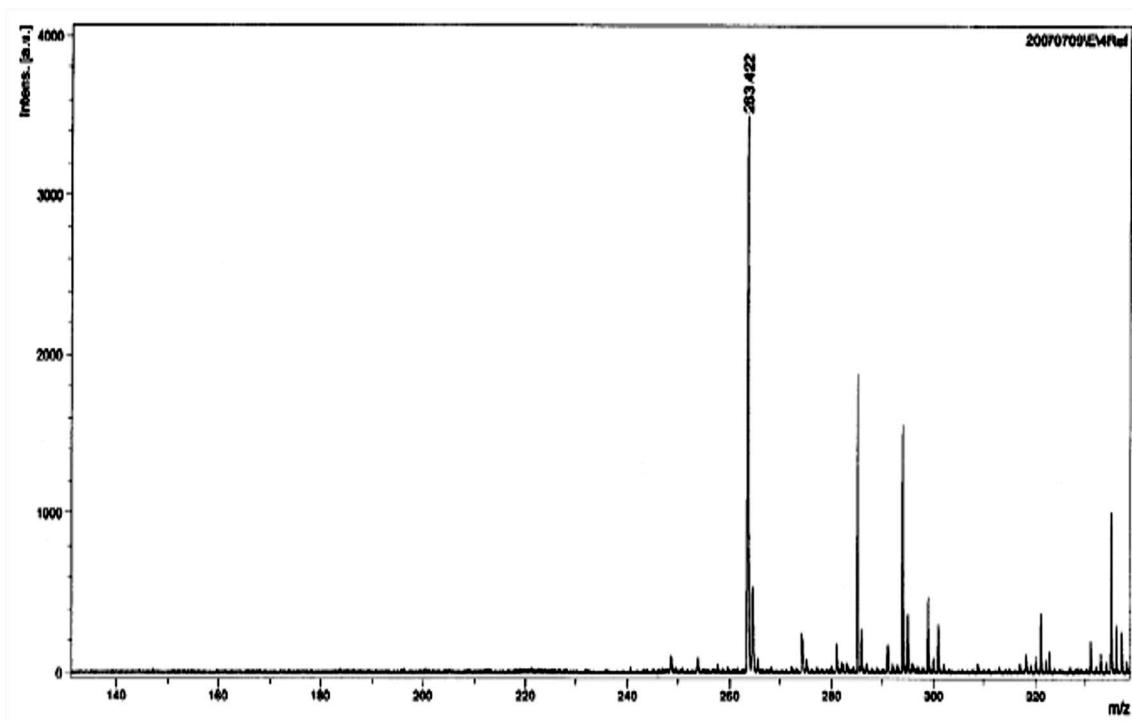


Figure S7. Mass spectrum of Methacr-L-Ser-NHBn.

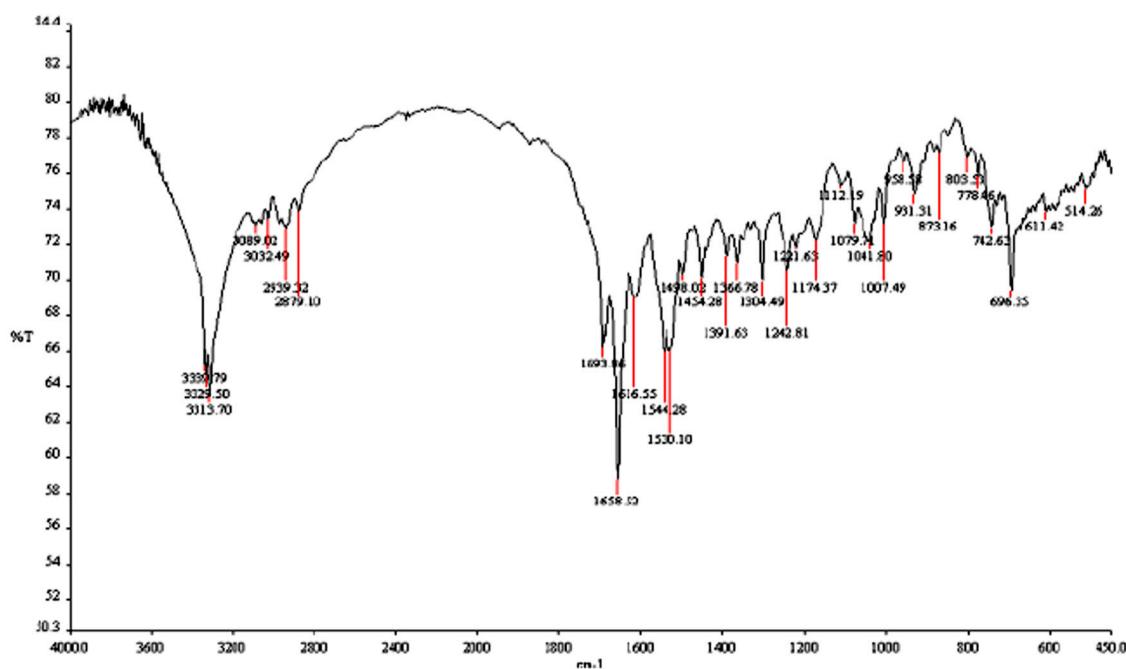


Figure S8. IR spectrum of Methacr-L-Ser-NHBn.