



## Supplementary data

## Exploring matrix effects on binding properties and characterization of cotinine molecularly imprinted polymer on paper-based scaffold

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**Figure S1.** Microscopic pictures of glass fiber membranes. (**a**) MIP paper-based scaffold (front view). (**b**) Bare glass fiber membrane (front view). (**c**) Stack of four bare glass fiber membranes (side view).



Figure S2. Chromatogram of standard mixture.



**Figure S3.** Infrared spectra of synthesized polymers. Red line represents MIP (nonextracted), blue line represents MIP (extracted), and black line represents NIP (focusing on 1720 cm<sup>-1</sup>).



\*+A; with agarose gel, +MIP; with MIP particles, +NIP; with NIP particles

**Figure S4.** Remaining cotinine standard after rebinding with different adsorbent materials. Black bar graph represents concentration of cotinine standard before rebinding.



Figure S5. Scatchard plot of NIP.



## Before MIP-paper based scaffold extraction

Figure S6. Chromatogram of eluted cotinine from MIP-paper based scaffold.

	Nicotine	Cotinine	Myosmine
Theoretical plate (N)	2480	11,142	9190
Capacity factor (k')	2.42	3.72	5.65
Selectivity $(\alpha)^a$	-	1.54	1.52
Resolution (R <sub>s</sub> ) <sup>a</sup>	-	5.27	6.22
HETP (µm)	111	22	27

**Table S1.** Chromatographic parameters for HPLC method.

<sup>a</sup> With respect to previous peak.

HETP, height equivalent to theoretical plate.

**Table S2.** Concentration of adsorbed cotinine ( $\mu$ g/mL) from MIP paper-based scaffold and bare paper-based scaffold (initial concentration of cotinine: 5  $\mu$ g/ mL).

Absorbent	Concentration of adsorbed	
	cotinine (µg/mL)	
Paper-based scaffold (with agarose; with MIPs)	$3.8 \pm 0.0810$	
Paper-based scaffold (with agarose; without MIPs)	$0.63 \pm 0.0042$	