

Supplementary Material
Enantioseparation in high performance liquid chromatography: preparation and evaluation of a vancomycin-based chiral stationary phase via surface-initiated atom transfer radical polymerization

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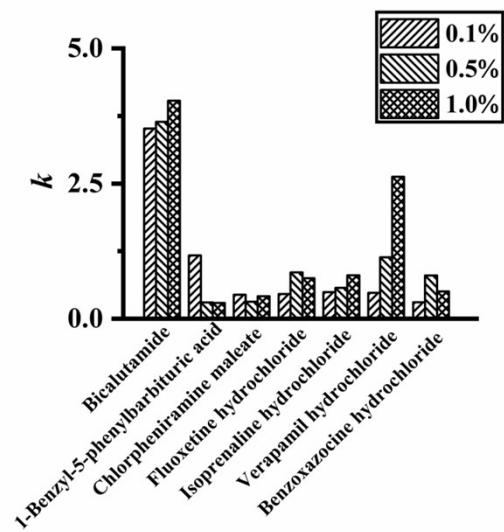


Fig. S1. Effect of TEA concentration in mobile phase on k of racemic drugs. Chromatographic conditions: mobile phase, TEAA solution (pH 3.5)/methanol (50/50, v/v); flow rate, 1 mL min⁻¹; injection volume, 5 μ L; detection temperature, 25 °C; detection wavelength, 224 nm.

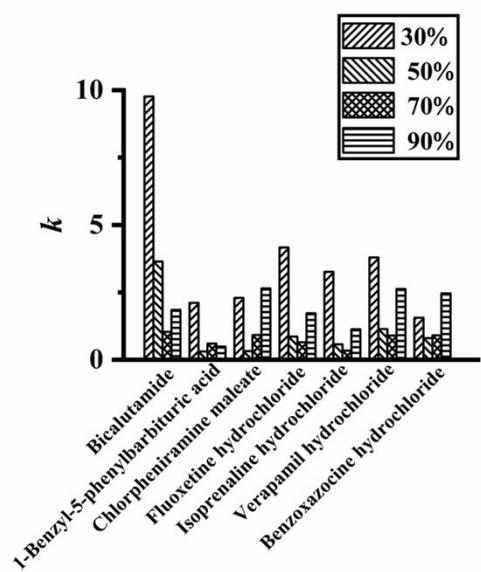


Fig. S2. Effect of methanol content in mobile phase on k of racemic drugs. Chromatographic conditions: mobile phase, 0.5% TEAA solution (pH 3.5)/methanol, flow rate, 1 mL min⁻¹; injection volume, 5 μ L; detection temperature, 25 °C; detection wavelength, 224 nm.

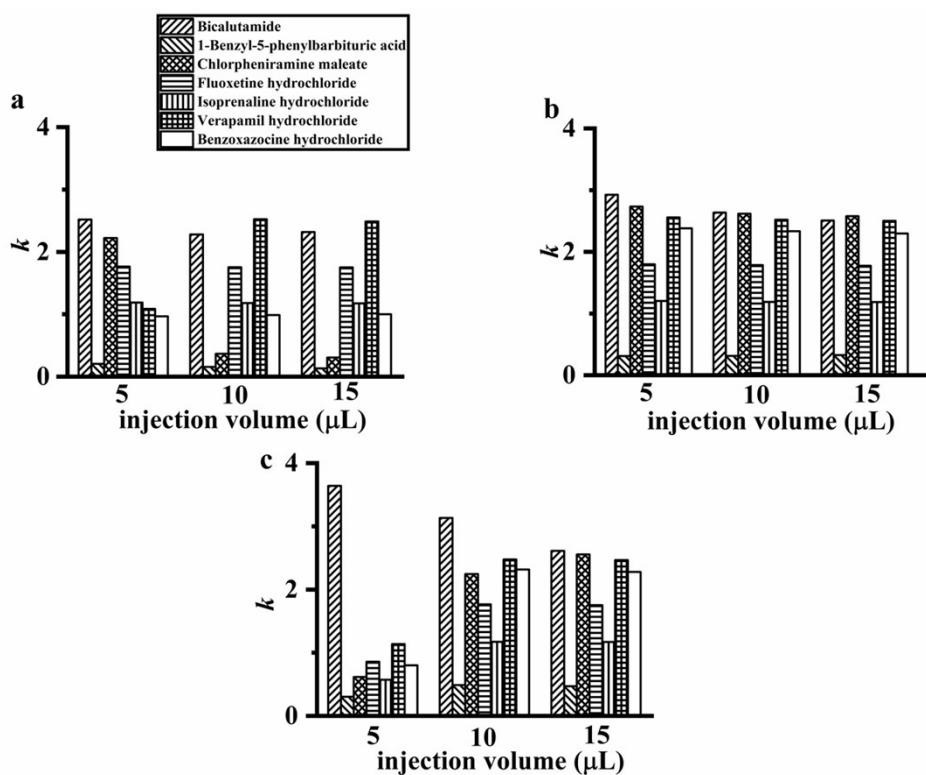


Fig. S3. Effect of injection volume on k of racemic drugs at the flow rate of (a) 0.6, (b) 0.8 and (c) 1 mL min^{-1} . Chromatographic conditions: mobile phase, 0.5% TEAA solution (pH 3.5)/methanol (50/50, v/v); detection temperature, 25 °C; detection wavelength, 224 nm.

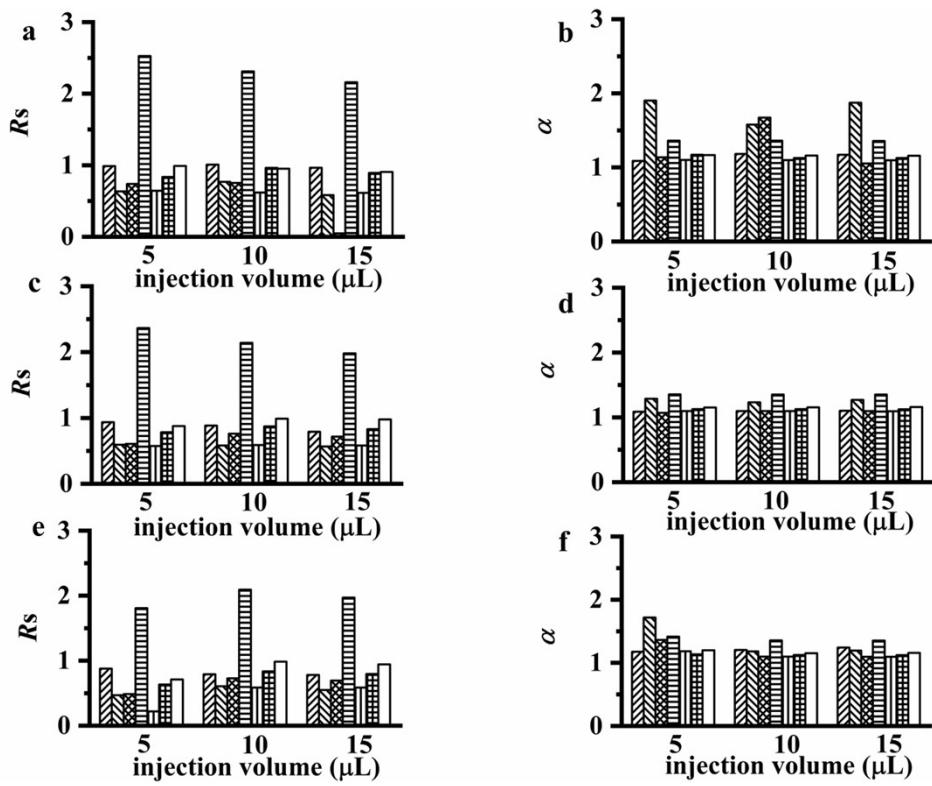


Fig. S4. Effect of injection volume on (a, c, e) R_s and (b, d, f) α of racemic drugs at the flow rate of (a, b) 0.6, (c, d) 0.8 and (e, f) 1.0 mL min^{-1} . Seven racemates as shown in Fig. S3.

Table S1 Values of ΔH^0 , ΔS^0 and ΔG^0 (25 °C) for benzoxazocine hydrochloride and chlorpheniramine maleate.

Analytes	ΔH^0	ΔS^0	ΔG^0	R
	(kJ mol ⁻¹)	(J mol ⁻¹ K ⁻¹)	(kJ mol ⁻¹)	
1-benzoxazocine hydrochloride	-2.682	-2.596	-2.682	0.9899
2-benzoxazocine hydrochloride	-2.587	-2.673	-2.586	0.9899
1-chlorpheniramine maleate	-2.702	-2.727	-2.702	0.9950
2-chlorpheniramine maleate	-2.362	-3.001	-2.361	0.9950

0.5% TEAA solution (pH 4.5)/methanol (50/50, v/v); flow rate, 1 mL min⁻¹; injection column, 5 µL; detection wavelength, 224 nm; column dimension, 150×4.6 mm I.D..