Supplementary Information

Figure S1. Comparison of preconcentration factor of studied parabens using CPE-DC193C and CPE-DC193C-βCD-IL.

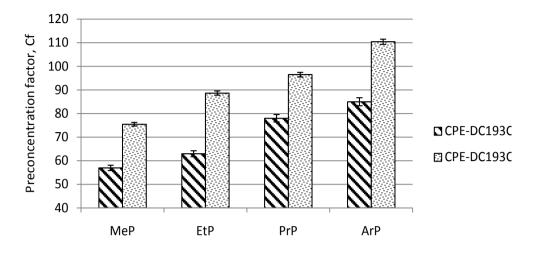
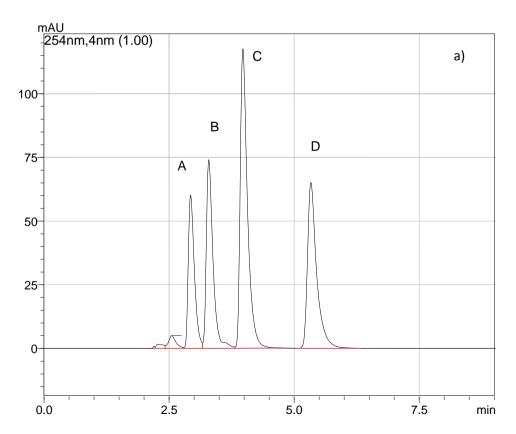


Figure S2. (a) Chromatogram of mixture of parabens; (b) Chromatogram of samples of sea water spiked with parabens. Amount of spiked at 400 ng/L of MeP, EtP, PrP and ArP. Cloud point conditions: 2.0 wt% (w/v) DC 193C. HPLC condition: Chromolith C18 column (100 mm × 4.6 mm). HPLC gradient conditions were used to separate the analytes using acetonitrile and deionized water, flow rate of 0.7 mL/min and detection at 254 nm. The gradient elution was performed as follows: 30% acetonitrile (0–2 min), ramped to 40% acetonitrile (3–5 min) and then ramped to 30% acetonitrile (5–8 min). A = methyl paraben, B = ethyl paraben, C = propyl paraben, D = benzyl paraben.



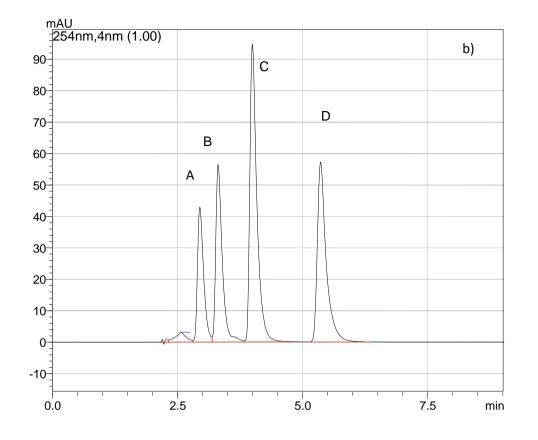
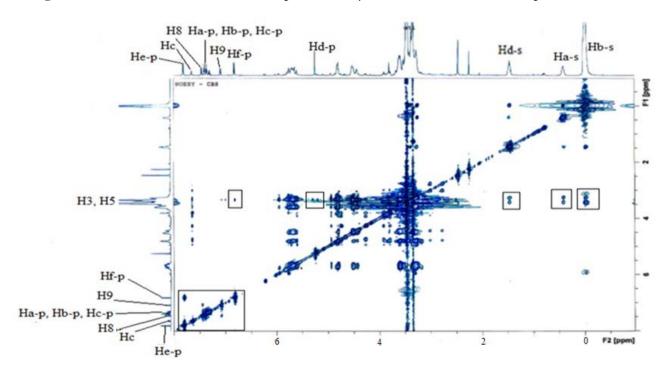


Figure S2. Cont.

Figure S3. Two-dimensional NOESY spectrum of β CD-IL-DC193C-ArP complex in DMSO-D₆.



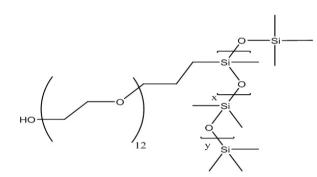


Figure S4. Structure of non-ionic surfactant DC193C.

Figure S5. Preparation of βCD functionalized IL (βCD-IL).

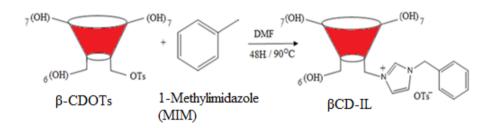


Figure S6. ¹H NMR spectrum of (**a**) β CD; and (**b**) β CD-OTs.

