

Supplementary Materials

Comparison of the Performance of Different Bile Salts in Enantioselective Separation of Palonosetron Stereoisomers by Micellar Electrokinetic Chromatography

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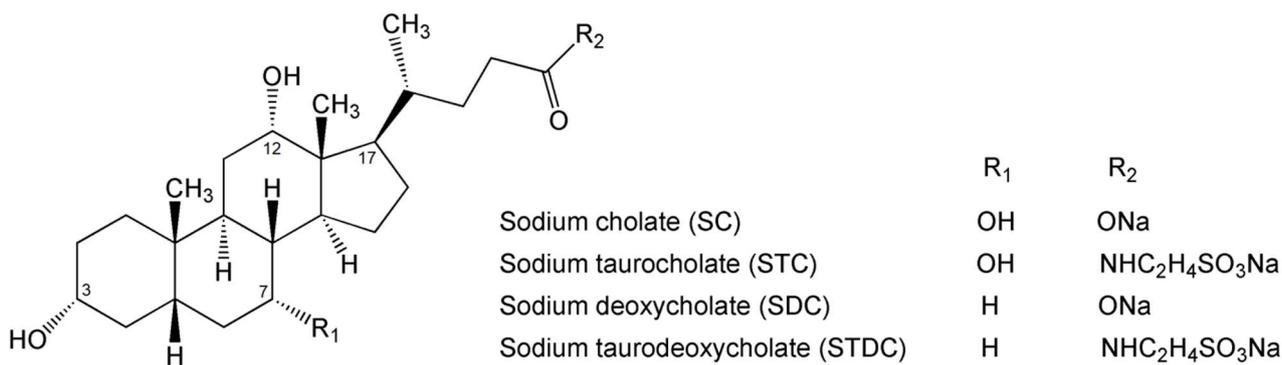


Figure S1. Molecular structures studied bile salts.

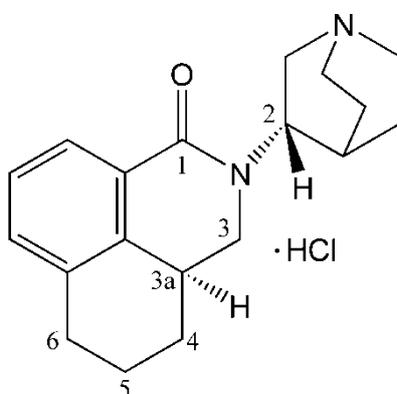


Figure S2. Molecular structure of palonosetron hydrochloride (PALO 3aS, 2S).

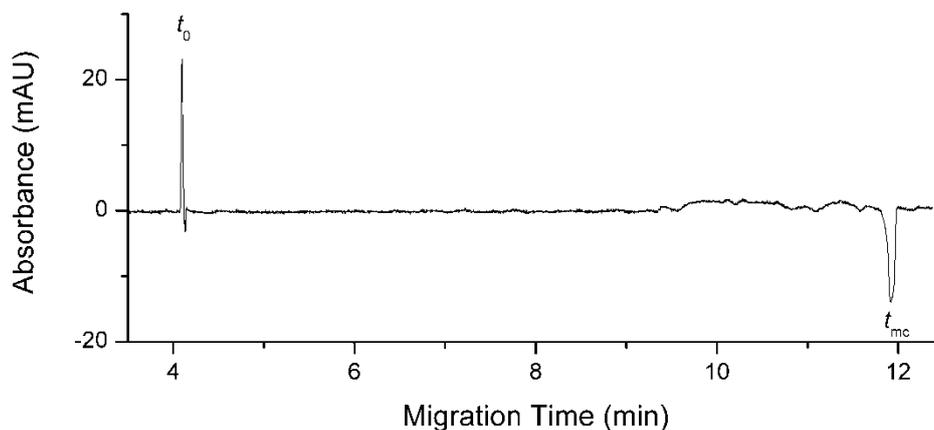


Figure S3. Determination of the migration time of micelles and the time window. Component of micellar solution (BGE) is 30 mM SDC in 30 mM sodium tetraborate buffer of pH 9.20. Capillary: id 50 μm , L_{tot} 60.0 cm, L_{eff} 50.0 cm. Capillary temperature: 20°C. Sample solution: 30 mM sodium tetraborate buffer of pH 9.20, containing 0.2 $\mu\text{L}\cdot\text{mL}^{-1}$ DMSO. Hydrodynamic injection at 10 kPa for 3 s. Detection wavelength: 205 nm. Applied voltage: 25 kV.

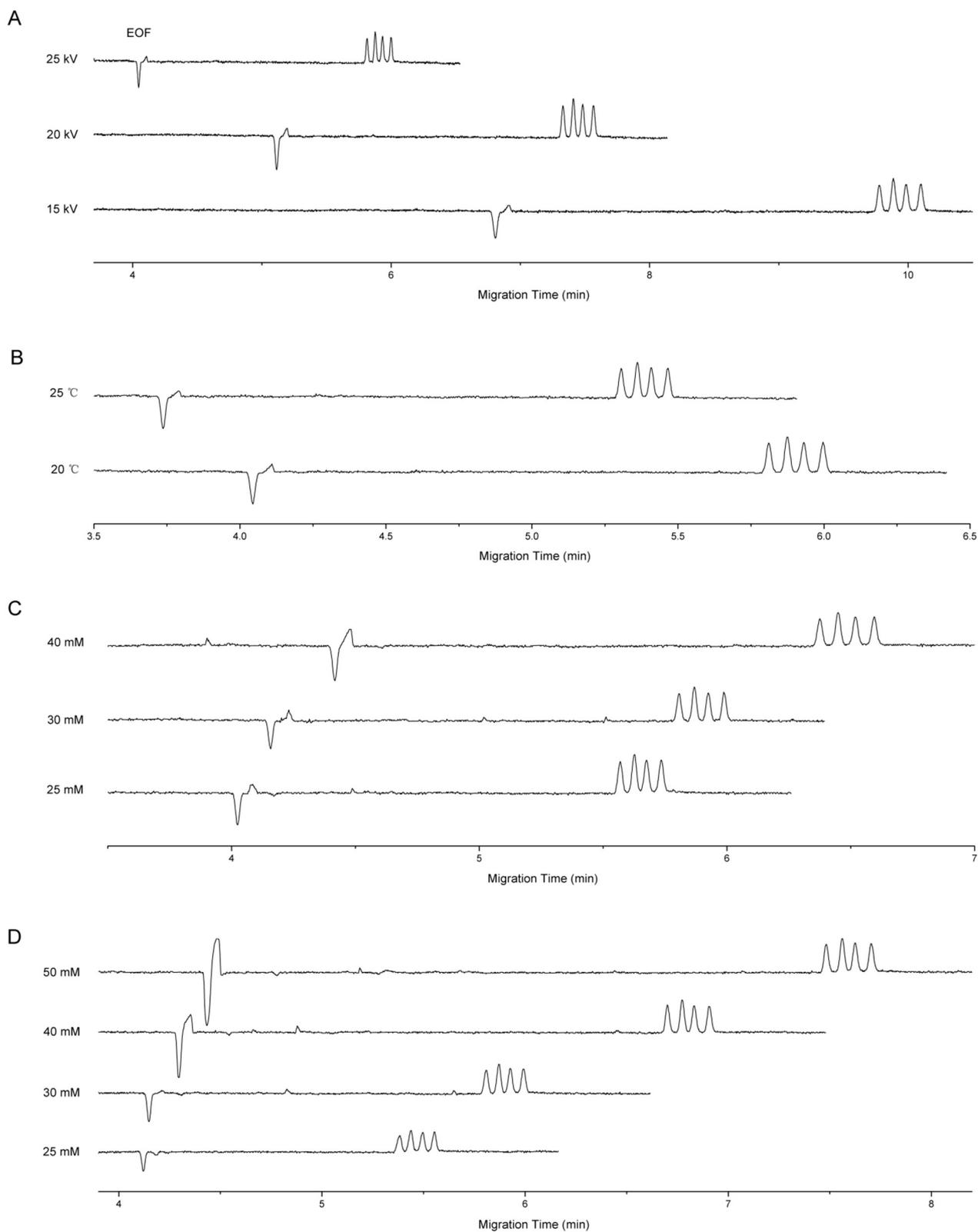


Figure S4. Electropherograms of PALO stereoisomers at different applied voltages (A), capillary temperatures (B), borate buffer concentrations (C) and STC surfactant

concentrations (D). Component BGE is 30 mM STC in 30 mM sodium tetraborate buffer of pH 9.20 except for specified. BGE Sample concentration: $0.1 \text{ mg} \cdot \text{mL}^{-1}$ for each stereoisomer. Hydrodynamic injection at 10 kPa for 1 s. Detection wavelength: 254 nm. Other CE conditions are the same as in Fig. S3 except for specified. The four peaks from left to right are in the order of PALO (3aS, 2S), PALO (3aR, 2R), PALO (3aS, 2R) to PALO (3aS, 2R) in all the electropherograms.