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Chiral Profiling Analysis of α -, β -, γ -, and δ -Blockers by Capillary Electrophoresis Using Dual Chiral Selectors

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Abstract

Simultaneous analysis of several β -blockers using a novel dual chiral selector system was achieved for their accurate chiral discrimination by chiral capillary electrophoresis (CE) in normal polarity mode. The CE system was operated using heptakis(2,6-di-*n*-methyl)- β -cyclodextrin (DM β -CD) dissolved in 100 mM phosphoric acid, triethanolamine buffer (pH 3) and (+)- or (-)-18-crown-6-tetracarboxylic acid (18-C-6-TA) as a co-chiral selector in DM β -CD solution. All eight investigated analytes were resolved using (+)-18-C-6-TA as a co-chiral selector in the presence of DM β -CD solution, whereas the other β -blockers, except propranolol, were resolved using (-)-18-C-6-TA as the second chiral selector in DM β -CD solution. Also, DM β -CD had a major effect on chiral discrimination compared with 18-C-6-TA as a co-chiral selector. Relative migration times to that of (S)-pindolol as an internal standard were characteristic of each enantiomer with good precision. The method showed good linearity with a correlation coefficient (r^2 , 0.993). The precision as a percentage of relative standard deviation (% RSD) and accuracy as a percentage of relative error (%)

RE) ranged from 0.5 to 9.6 and from 5.7 to 9.1, respectively. These were adequate for the chiral assay of the α -blockers investigated. Thus, the present chiral profiling method is expected to be useful for accurate chiral discrimination and optical purity tests of chiral α -blockers and their related drugs.

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