European Pharmacopoeia method



Timolol maleate: Enantiomeric purity

Column : CHIRALCEL[®] OD-H 0.46cmΦ × 25cmL

Mobile phase : Diethylamine R / 2-Propanol R / Hexane R = 2 / 40 / 960 (v / v / v)

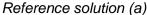
Flow rate : 1.0mL/min.

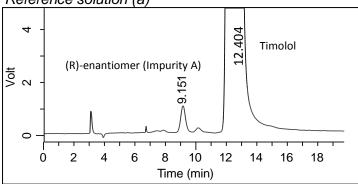
Injection volume : 5µL UV detection : 297nm

System suitability

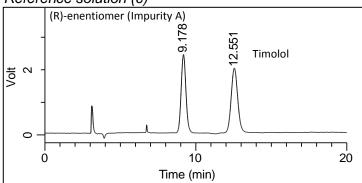
Reference solution (a):Dissolve 30.0 mg of *Timolol maleate CRS* in the solvent mixture and dilute to 10 mL with the solvent mixture.

Reference solution (c):Dilute 1 mL of reference solution (a) to 100 mL with the solvent mixture. Mix 1 mL of this solution with 1 mL of reference solution (b).





Reference solution (c)



	Requirement	Result
Resolution	Minimum 4.0 between the peaks due to impurity A and the (S)-enantiomer (reference solution (c))	4.9
Retention time	The retention times of the principal peaks due to the (S) -enantiomer in the chromatograms obtained with the test solution and reference solution (a) are identical.	Identical

For details of monograph, please check pharmacopoeia.