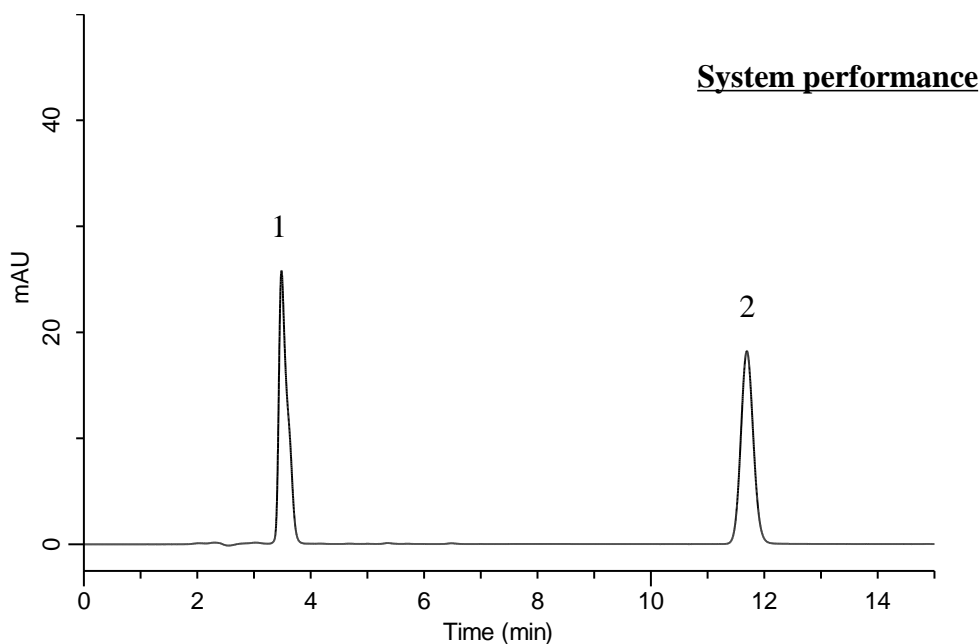


Analysis of Hydrochlorothiazide

(Under the Condition of the Japanese Pharmacopoeia,
Candesartan cilexetil and Hydrochlorothiazide Tablets)



Conditions

System : GL7700 HPLC system
Column : Inertsil ODS-4
 (4 μ m, 150 x 4.6 mm I.D.)
Column Cat. No. : 5020-89608
Eluent : A) CH₃CN
 B) Buffer*
 A/B = 11/9, v/v
Flow Rate : 0.57 mL/min
Col. Temp. : 25 °C
Detection : UV 254 nm (UV7750 UV Detector)
Injection Vol. : 10 μ L
Sample : Standard

Analyte:

1. Hydrochlorothiazide 3.8 mg/L
 2. Candesartan cilexetil 2.5 mg/L

Resolution (1, 2) : 24.2 (≥ 7)

*Dissolve 7.80 g of sodium dihydrogenphosphate dihydrate in 900 mL of water.
 Adjust pH 5.5 by sodium hydroxide.
 Add water to make 1,000 mL.