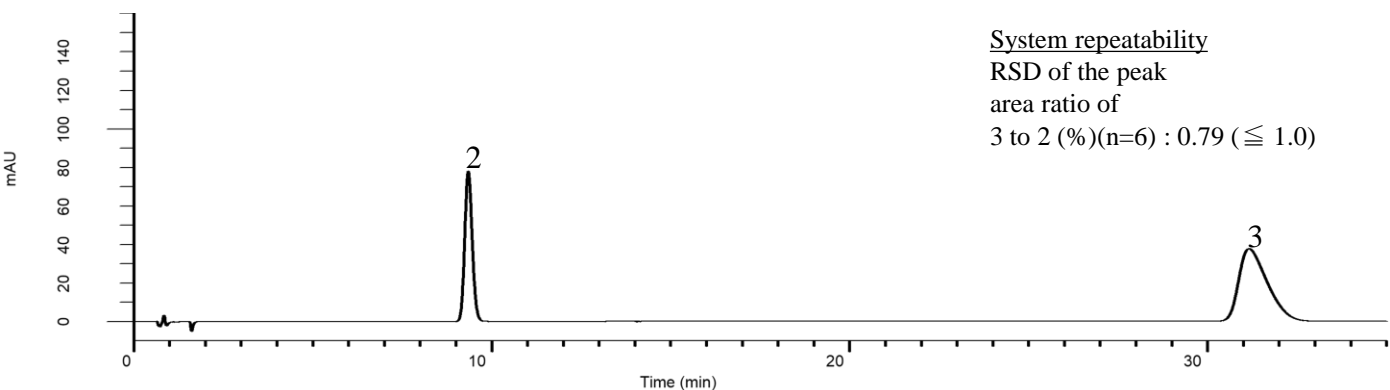
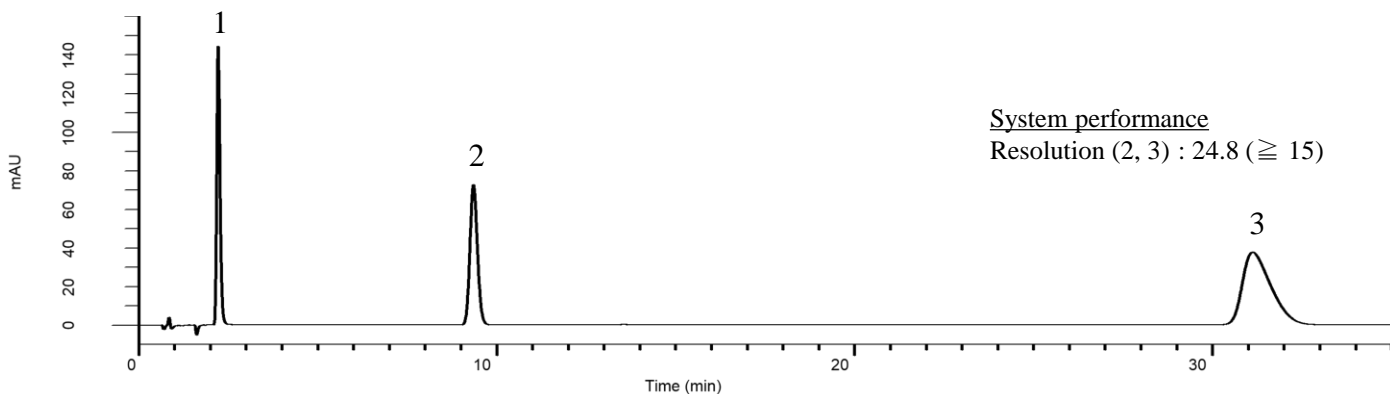


Analysis of Candesartan cilexetil

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



Conditions

System : GL-7400 HPLC system
Column : Inertsil ODS-4
 (5 μ m, 150 x 3.9 mm I.D.)
Column Cat. No. : 5020-87023
Eluent : A) CH₃CN
 B) 0.7 % Triethylamine in H₂O (pH 6.5, H₃PO₄)
 A/B = 80/50, v/v
Flow Rate : 1.55 mL/min
Col. Temp. : 25 °C
Detection : UV 238 nm (GL-7452 PDA Detector)
Injection Vol. : 10 μ L
Sample : Standard

Analyte:

1. Amlodipine besylate	70 mg/L
2. <i>p</i> -Hydroxybenzoic acid <i>n</i> -butyl ester	80 mg/L
3. Candesartan cilexetil	160 mg/L