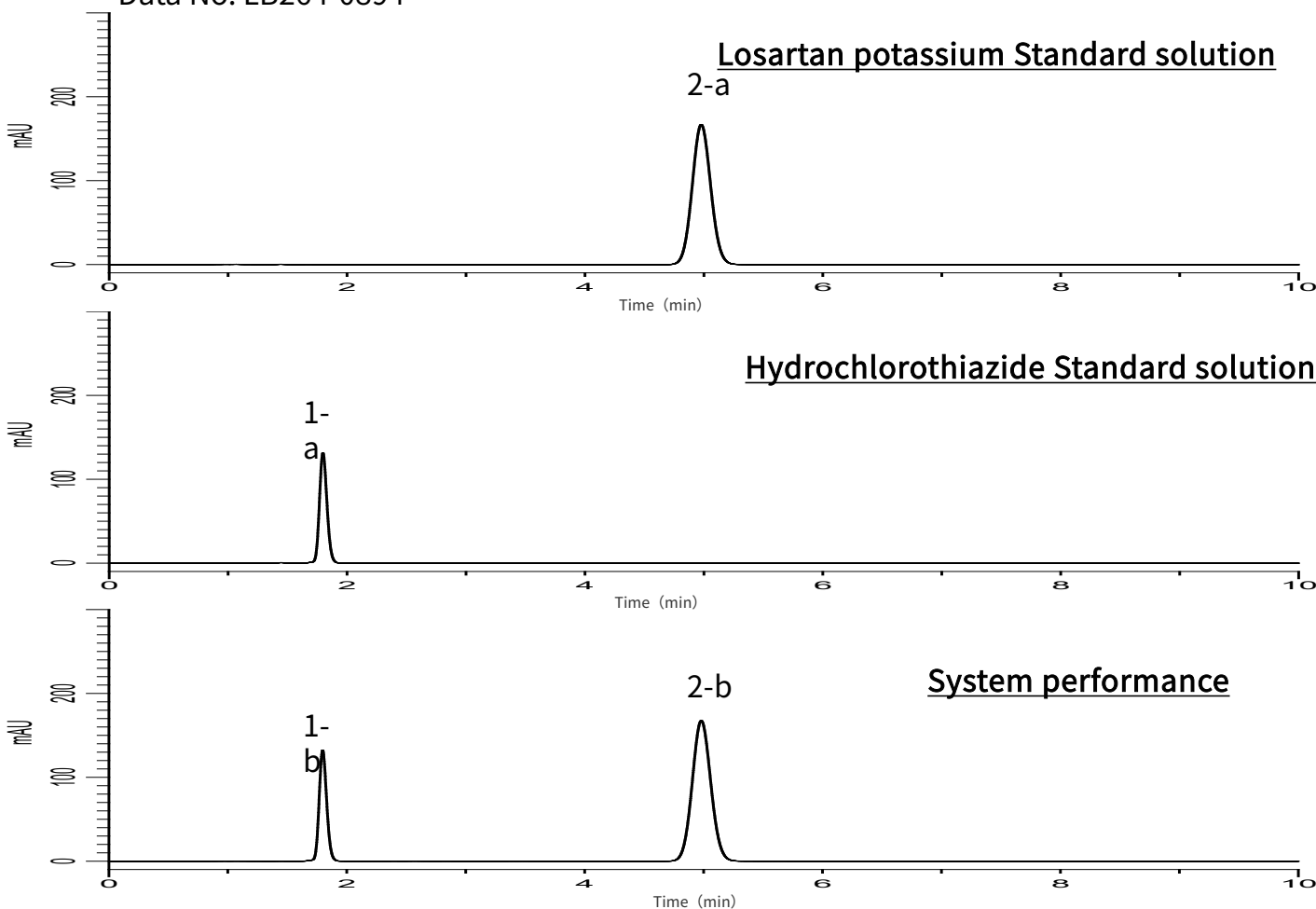


Analysis of Losartan potassium and Hydrochlorothiazide (Under the Condition of the Japanese

Pharmacopoeia,

Losartan potassium and Hydrochlorothiazide Tablets)

Data No. LB264-0894



Conditions

System : GL7700 HPLC system
Column : Inertsil C8-3 (10 μ m, 250 x 4.6 mm I.D.)
Column Cat. No. : 5020-01642
Eluent : A) CH₃CN
 : B) 10mM KH₂PO₄ (pH 2.5, H₃PO₄)
 : A/B = 40/60, v/v
Flow rate : 2.25 mL/min
Col. Temp. : 35 °C
Detection : UV 230 nm (UV7750 UV Detecor)
Injection Vol. : 20 μ L
Sample : Standard

Analyte:

1. Hydrochlorothiazide 14 mg/L
 2. Losartan potassium 55 mg/L

Resolution (1-b, 2-b) : 15.7 (≥ 10)
 RSD of the peak
 area of 1-a (%) (n=6) : 0.07 (≤ 1.0)
 RSD of the peak
 area of 2-a (%) (n=6) : 0.04 (≤ 1.0)