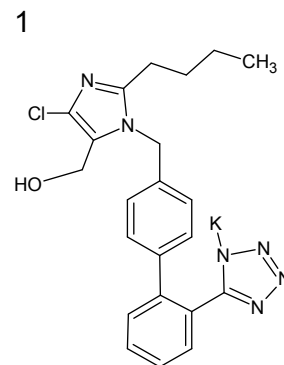
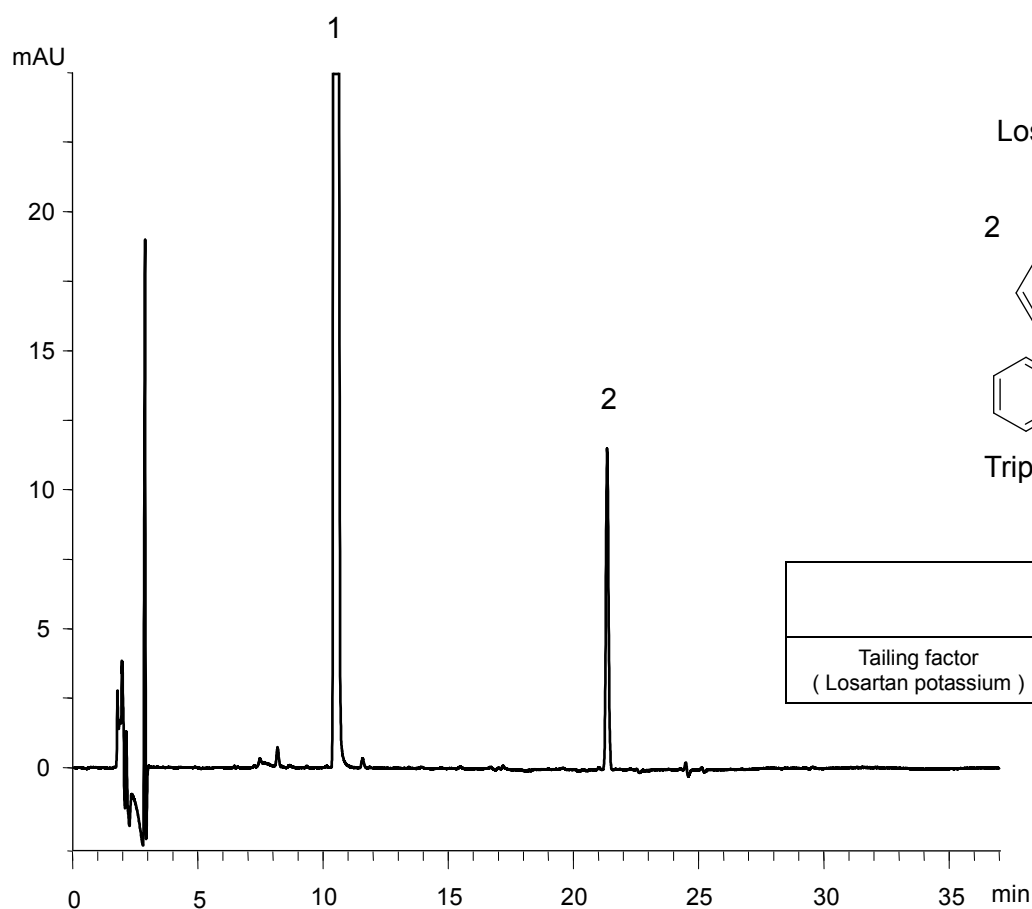


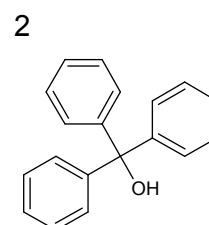
ロサルタンカリウム (米国薬局方記載条件)
Losartan potassium (The United States Pharmacopeia)

D111122A

System suitability solution *
(0.3 mg/mL Losartan potassium, 0.002 mg/mL Triphenylmethanol)



Losartan potassium



Triphenylmethanol

	System suitability requirement	Result
Tailing factor (Losartan potassium)	≤ 1.6	1.16

Column : YMC-Triart C18 (5 μm, 12 nm)
250 X 4.0 mmI.D.

Eluent : A) 0.1% phosphoric acid
B) acetonitrile
25-90%B (0-25 min), 90%B (25-35 min)

Flow rate : 1.0 mL/min

Temperature : 25°C

Detection : UV at 220 nm

Injection : 10 μL

(The United States Pharmacopeia 34th; Chromatographic purity)

* System suitability solution was prepared from Losartan potassium supplied as a reagent for laboratory use.