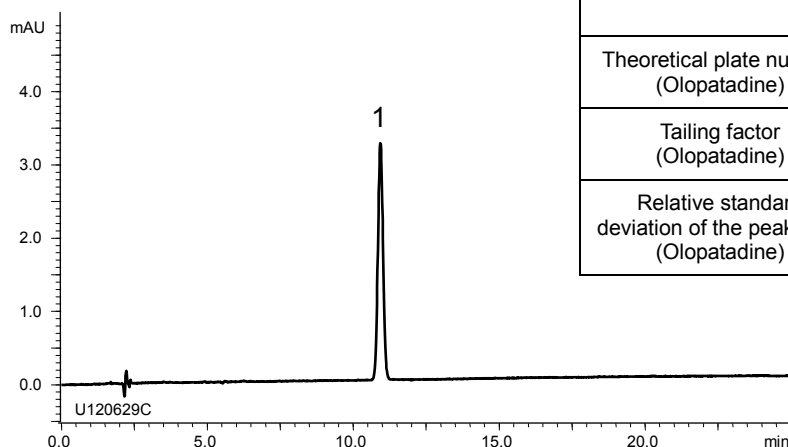


オロパタジン塩酸塩 (日本薬局方収載原案記載条件)
Olopatadine hydrochloride (The draft for the Japanese Pharmacopoeia)

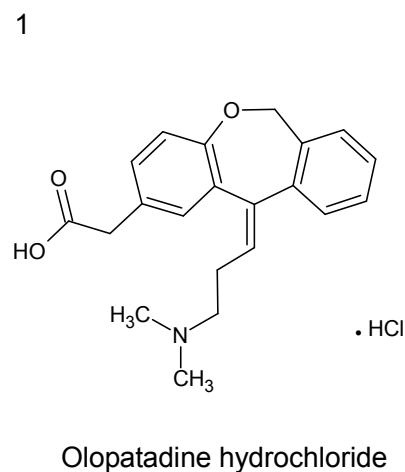
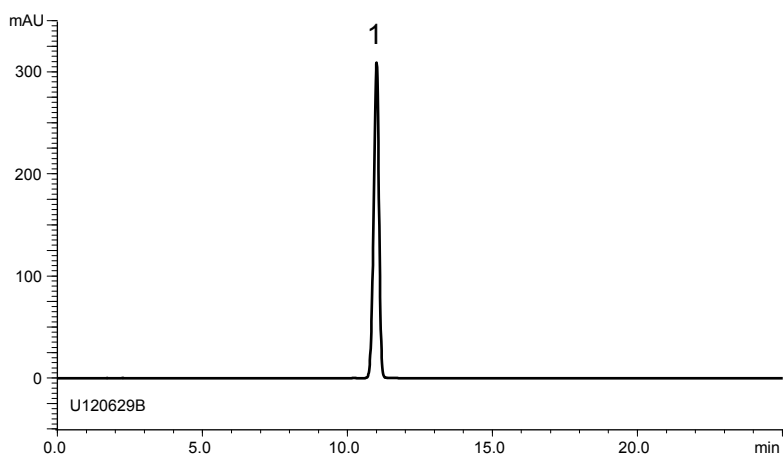
U120724A

A) Standard solution*¹
(0.005 mg/mL Olopatadine HCl)



	System suitability requirement	Result
Theoretical plate number (Olopatadine)	≥ 8000	18100
Tailing factor (Olopatadine)	≤ 2.0	1.08
Relative standard deviation of the peak area (Olopatadine)	$\leq 1.0\%$	0.07%

B) Sample solution*¹
(0.5 mg/mL Olopatadine HCl)



Column : YMC-Triart C8 (5 μ m, 12 nm)
250 X 4.6 mmI.D.

Eluent : phosphate buffer (pH 3.5)*²/acetonitrile (11/9) containing 8 mM sodium lauryl sulfate
*² Dissolve 8.6 g of KH_2PO_4 in 1000 mL of water, adjust pH 3.5 with H_3PO_4 (49→10000)

Flow rate : 1.1 mL/min (adjust the flow rate so that the retention time of olopatadine is about 11 min)

Temperature : 40°C

Detection : UV at 299 nm

Injection : 20 μ L

(The draft for the Japanese Pharmacopoeia; Related substances)

*¹ All standard and sample solutions were prepared from Olopatadine hydrochloride supplied as a reagent for laboratory use.