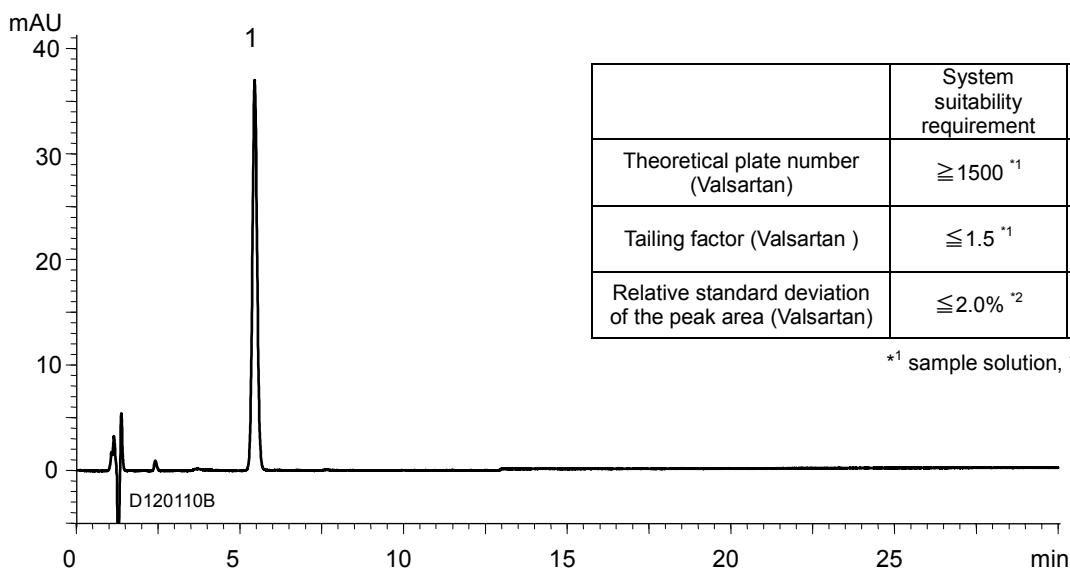


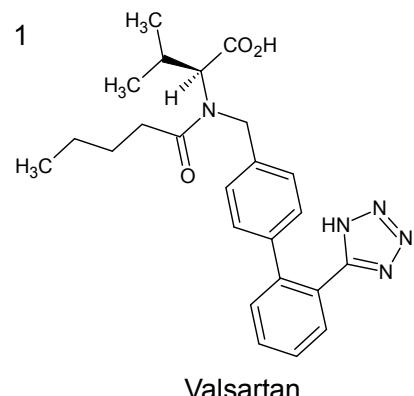
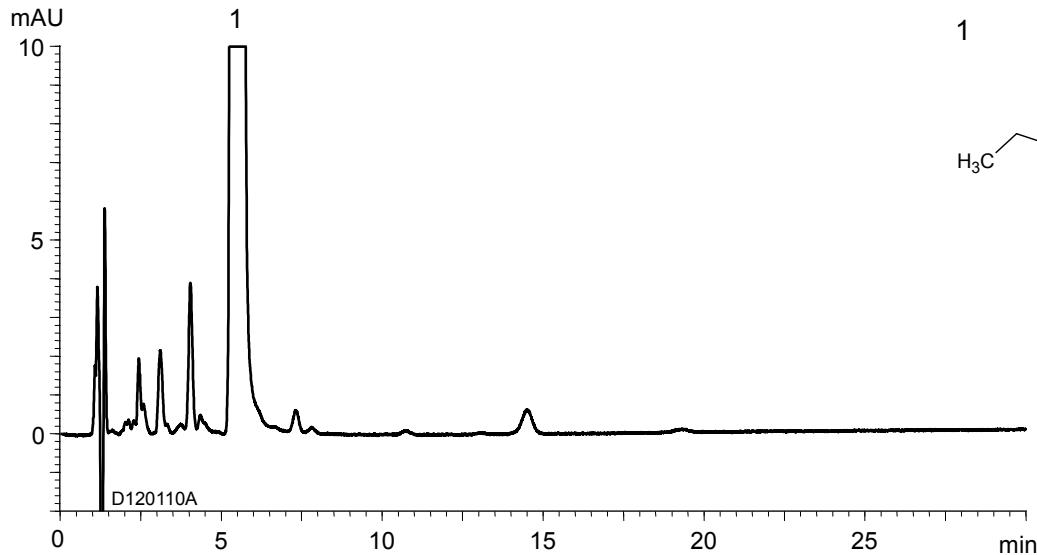
バルサルタン（日本薬局方収載原案記載条件）

Valsartan (The draft for the Japanese Pharmacopoeia)

D120110D

A) Standard solution^{*1} (0.005 mg/mL Valsartan)

	System suitability requirement	Result for standard solution	Result for sample solution
Theoretical plate number (Valsartan)	≥ 1500 ^{*1}	6700	N.C.
Tailing factor (Valsartan)	≤ 1.5 ^{*1}	1.06	N.C.
Relative standard deviation of the peak area (Valsartan)	$\leq 2.0\%$ ^{*2}	0.10%	—

^{*1} sample solution, ^{*2} standard solutionB) Sample solution^{*1} (0.5 mg/mL Valsartan)

Column	: YMC-Triart C18 (5 μ m, 12 nm) 125 X 3.0 mmI.D.
Eluent	: acetonitrile/water/acetic acid (50/50/0.1)
Flow rate	: 0.45 mL/min (<i>adjust the flow rate so that the retention time of valsartan is about 5 min</i>)
Temperature	: 25°C
Detection	: UV at 225 nm
Injection	: 10 μ L

(The draft for the Japanese Pharmacopoeia; Related substances)

^{*1} Standard and sample solutions were prepared from Valsartan supplied as a reagent for laboratory use.