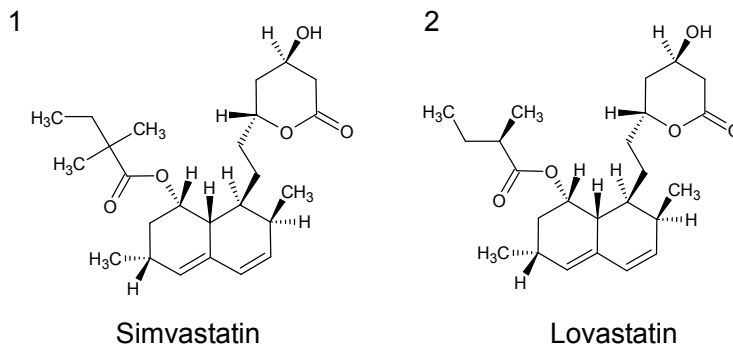
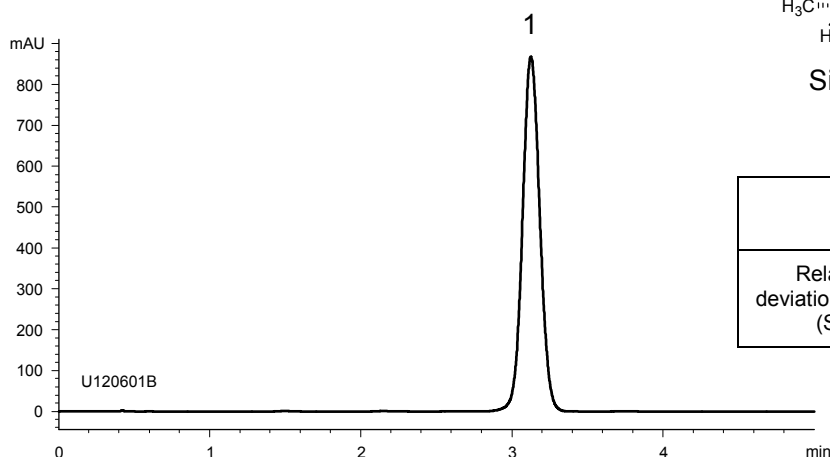


シンバスタチン（日本薬局方記載条件）
Simvastatin (The Japanese Pharmacopoeia)

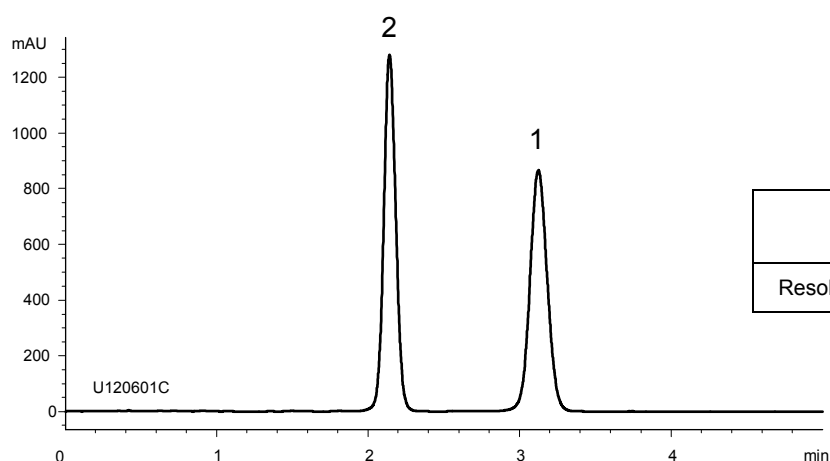
U120619B

A) Standard solution*
(1.5 mg/mL Simvastatin)



	System suitability requirement	Result
Relative standard deviation of the peak area (Simvastatin)	≤ 1.0%	0.44%

B) System suitability solution*
(1.5 mg/mL Lovastatin, 1.5 mg/mL Simvastatin)



	System suitability requirement	Result
Resolution (1, 2)	≥ 3	5.6

Column : YMC-Triart C18 (3 μm, 12 nm)
35 X 4.6 mm I.D.

Eluent : acetonitrile/water/phosphoric acid (50/50/0.05)

Flow rate : 3.0 mL/min (*adjust the flow rate so that the retention time of simvastatin is about 3 min*)

Temperature : 25°C

Detection : UV at 238 nm

Injection : 5 μL

(The Japanese Pharmacopoeia 16th; Assay)

* All solutions were prepared from Simvastatin supplied as a reagent for laboratory use.