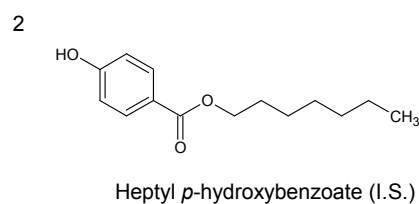
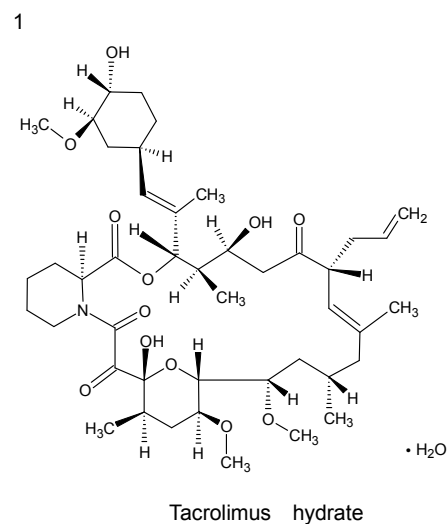
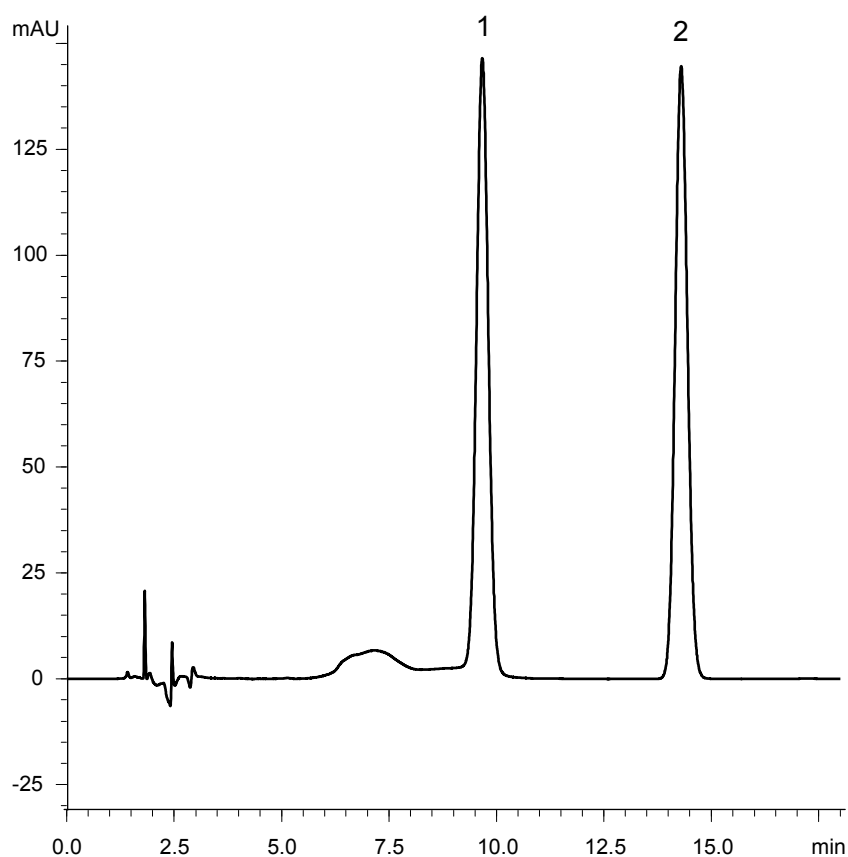


タクロリムス水和物（日本薬局方記載条件）  
Tacrolimus hydrate (The Japanese Pharmacopoeia)

U120322D

Sample solution\*  
(0.5 mg/mL Tacrolimus, 0.15 mg/mL Heptyl *p*-hydroxybenzoate)



	System suitability requirement	Result
Resolution (1, 2)	$\geq 6$	8.8
Relative standard deviation of the peak area ratio of 1 to 2	$\leq 1.0\%$	0.93%

Column : YMC-Triart C18 (5  $\mu$ m, 12 nm)  
150 X 4.6 mm I.D.  
Eluent : 2-propanol/THF/water (2/2/5)  
Flow rate : 0.7 mL/min (*adjust the flow rate so that the retention time of tacrolimus is about 10 min*)  
Temperature : 50°C  
Detection : UV at 220 nm  
Injection : 10  $\mu$ L  
(The Japanese Pharmacopoeia 16th; Assay)

\*Sample solution was prepared from Tacrolimus supplied as a reagent for laboratory use.