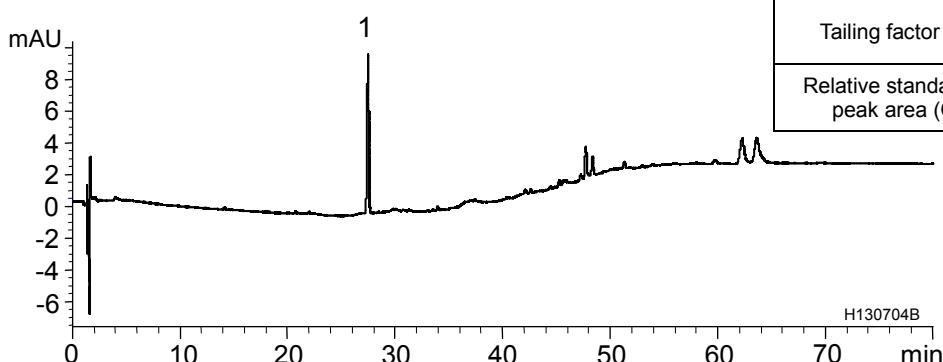


クロピドグレル硫酸塩（日本薬局方原案記載条件）

Clopidogrel Sulfate (The draft for the Japanese Pharmacopoeia) H130704D

(A) Standard solution<sup>\*1</sup>

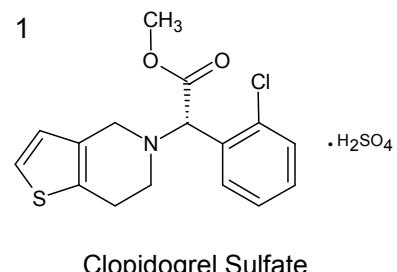
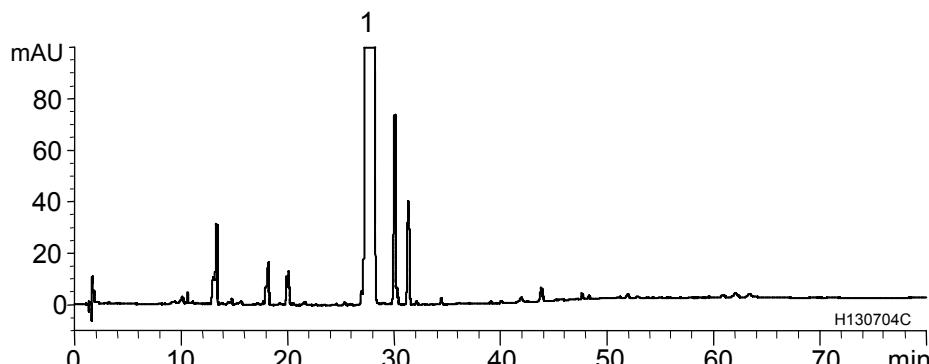
(6.5 µg/mL Clopidogrel sulfate)



	System suitability requirement	Result
Theoretical plate number (Clopidogrel)	≥60000	155300
Tailing factor (Clopidogrel)	≤2.0	0.97
Relative standard deviation of peak area (Clopidogrel)	≤2.0%	0.30

(B) Sample solution<sup>\*1</sup>

(6.5 mg/mL Clopidogrel sulfate)



Column	: YMC-Pack Pro C18 (5 µm, 12 nm) 150 X 4.0 mmI.D.
Eluent	: A) buffer <sup>*2</sup> /methanol (19/1) B) acetonitrile/methanol (19/1) <sup>*2</sup> Dissolve 0.87 g of sodium 1-pentanesulfonate in 1000 mL water, adjust pH 2.5 with H <sub>3</sub> PO <sub>4</sub> 10.5%B (0-3 min), 10.5-68.5%B (3-48 min), 68.5%B (48-68 min)
Flow rate	: 1.0 mL/min
Temperature	: 30°C
Detection	: UV at 220 nm
Injection	: 10 µL
(The draft for the Japanese Pharmacopoeia; Related substances)	

<sup>\*1</sup> All standard and sample solutions were prepared from Clopidogrel sulfate supplied as a reagent for laboratory use.