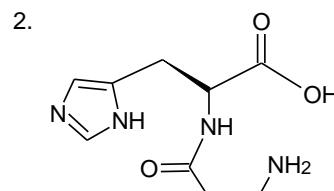
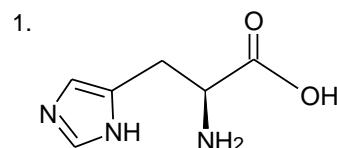
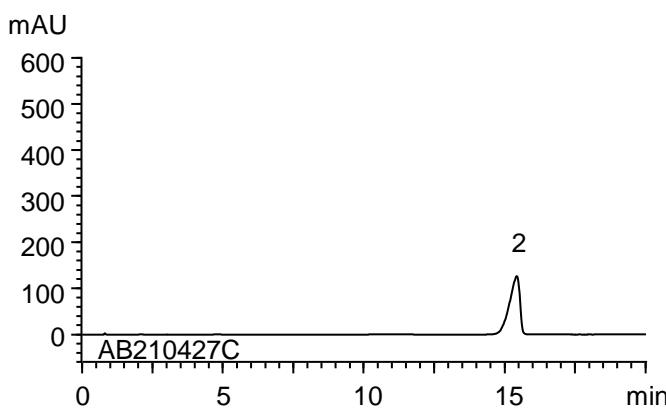
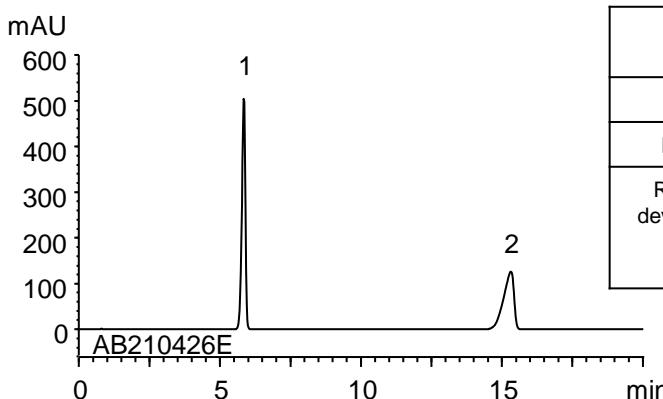


ポラプレジンク（日本薬局方記載条件）

Polaprezinc (The Japanese Pharmacopoeia)

AB210510A

(A) Standard solution^{*2} (0.2 mg/mL L-Carnosine)(B) System performance test solution^{*2}
(0.25 mg/mL L-Histidine, 0.2 mg/mL L-Carnosine)

	System suitability requirement	Result
Elution order	1, 2	1, 2
Resolution (1,2)	≥ 12	19
Relative standard deviation of the peak area (n=6) (L-Carnosine)	$\leq 1.0\%$	0.2%

- Column : YMC-Triart C18 (5 μm , 12 nm)
150 X 4.6 mmI.D.
- Eluent : phosphate buffer (pH 3.5)^{*1} containing 2.22 g/L sodium 1-octanesulfonate /acetonitrile (90/10)
- *1 Dissolve 1.4 g of KH_2PO_4 in 1000 mL of water, adjust pH 3.5 with 1% H_3PO_4 .
- Flow rate : 1.35 mL/min (adjust the flow rate so that the retention time of L-Carnosine is about 15 min)
- Temperature : 45°C
- Detection : UV at 210 nm
- Injection : 10 μL

(The Japanese Pharmacopoeia 17th 2nd supplement; Assay (1) Polaprezinc)

^{*2}All system performance test and standard solutions were prepared from L-Carnosine supplied as a reagent for laboratory use.