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L-Proline		
Issued Date: Dec, 4, 2015		

## L-Proline<sup>1</sup>

C<sub>5</sub>H<sub>9</sub>NO<sub>2</sub>: 115.13

L-Proline contains not less than 99.0 percent and not more than 101.0 percent of L-Proline (C<sub>5</sub>H<sub>9</sub>NO<sub>2</sub>), calculated on the dried basis.

### Description

White crystals or white crystalline powder; slightly sweet taste.

Very soluble in water and in formic acid, and slightly soluble in ethanol (99.5).

It is deliquescent.

### Identification

Compare the infrared absorption spectrum of the sample with that of the standard by potassium bromide disc method.

### Specifications

Item	Limit	Test
Specific rotation [ $\alpha$ ] <sub>D</sub> <sup>20</sup>	-84.5 to -86.0°	AJI TEST 1 [Calculated on the dried basis, C=4, H <sub>2</sub> O] <sup>2</sup>
State of solution (Transmittance)	Clear and colorless Not less than 98.0%	AJI TEST 2 [1.0g in 10mL of H <sub>2</sub> O, spectrophotometer, 430nm, 10mm cell thickness]
Chloride (Cl)	Not more than 0.020%	AJI TEST 3 [0.5g, A-1, ref: 0.28mL of 0.01mol/L HCl]
Ammonium (NH <sub>4</sub> )	Not more than 0.02%	AJI TEST 4 [D-1]
Sulfate (SO <sub>4</sub> )	Not more than 0.020%	AJI TEST 5 [0.85g, (1), ref: 0.35mL of 0.005mol/L H <sub>2</sub> SO <sub>4</sub> ]
Iron (Fe)	Not more than 10ppm	AJI TEST 6 [1.0g, A-1, ref: 1.0mL of Iron Std. (0.01mg/mL)]
Heavy metals (Pb)	Not more than 10ppm	AJI TEST 7 [1.0g, (1), ref: 1.0mL of Pb Std. (0.01mg/mL)]
Arsenic (As <sub>2</sub> O <sub>3</sub> )	Not more than 1ppm	AJI TEST 8 [2.0g, (1), ref: 2.0mL of As <sub>2</sub> O <sub>3</sub> Std.]
Related substances	1)Conforms <sup>3</sup>  2)Any unspecified impurity Not more than 0.20% Total impurities Not more than 0.50%	AJI TEST 9 [Test sample: 50μg, B-6-a, control; L-Pro 0.25μg] AJI TEST 26 <sup>4</sup>
Loss on drying	Not more than 0.30%	AJI TEST 11 [1g, at 105°C for 3 hours]
Residue on ignition (Sulfated)	Not more than 0.10%	AJI TEST 13 [1g, at 550°C to 650°C for 3 hours]

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### Specifications (cont'd)

Item	Limit	Test
Assay	99.0 to 101.0%	AJI TEST 14 [Calculated on the dried basis, 120mg, (1), 3mL of formic acid, 50mL of acetic acid (100), 0.1mol/L HClO <sub>4</sub> 1mL=11.51mg C <sub>5</sub> H <sub>9</sub> NO <sub>2</sub> ]
pH	5.9 to 6.9	AJI TEST 33 [1.0g in 10mL of H <sub>2</sub> O]

The test for Endotoxin when the material will be used for manufacturing parenteral products is as follows:

Item	Limit	Test
Endotoxin	Less than 6.0EU/g	AJI TEST 34 [C=1, kinetic-turbidimetric technique]

<sup>1</sup> This product, in terms of actual quality, conforms to USP, EP, and JP.

<sup>2</sup> Temperature coefficient of  $[\alpha]_D^{25}$ : 0.0°

<sup>3</sup> Any secondary spot in the chromatogram obtained from the Test Solution is less intense than the principal spot in the chromatogram obtained from the Standard Solution: the number of those spots is not more than four and not more than 2.0% of the total impurities found.

<sup>4</sup> Disregard limit: 0.05%

**End of document**