Amino Acids Specifications / Monographs		1/2
L-Threonine		
Issued Date: Dec, 4, 2015		

L-Threonine¹

C₄H₉NO₃: 119.12

L-Threonine, when dried, contains not less than 99.0 percent and not more than 101.0 percent of L-Threonine ($C_4H_9NO_3$).

Description

White crystals or crystalline powder; slightly sweet taste.

Freely soluble in formic acid, soluble in water, practically insoluble in ethanol (95).

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 $[\alpha]_D^{20}$	-27.6 to -29.0°	AJI TEST 1
		[Dried sample, $C=6$, H_2O] ²
State of solution	Clear and colorless	AJITEST 2
(Transmittance)	Not less than 98.0%	[1.0g in 10mL of H ₂ 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 ₄)	Not more than 0.02%	AJI TEST 4
		[D-1]
Sulfate (SO ₄)	Not more than 0.020%	AJI TEST 5
		[0.85g, (1), ref: 0.35mL of 0.005mol/L H ₂ SO ₄]
Iron (Fe)	Not more than 10ppm	AJI TEST 6
		[0.75g, B-1, ref: 0.75mL of Iron Std. (0.01mg/mL)]
Heavy metals (Pb)	Not more than 10ppm	AJITEST 7
		[2.0g, (1), ref: 2.0mL of Pb Std. (0.01mg/mL)]
Arsenic (As ₂ O ₃)	Not more than 1ppm	AJI TEST 8
		[2.0g, (1), ref: 2.0mL of As ₂ O ₃ Std.]
Related substances	1)Conforms ³	AJITEST 9
		[Test sample: 50µg, B-6-a, control; L-Thr 0.25µg]
	2)Any unspecified impurity	AJI TEST 26 ⁴
	Not more than 0.20%	
	Total impurities	
Τ 1 '	Not more than 0.50%	A HATTE CITE 11
Loss on drying	Not more than 0.20%	AJITEST 11
D 11 1 12	NT / 4 0400/	[1g, at 105°C for 3 hours]
Residue on ignition	Not more than 0.10%	AJITEST 13
(Sulfated)		[1g, at 550°C to 650°C for 3 hours]

AJINOMOTO CO,. INC.

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Specifications (cont'd)

Item	Limit	Test
Assay	99.0 to 101.0%	AJI TEST 14
		[Dried sample, 120mg, (1), 3mLof formic acid, 50mL of acetic
		acid (100), 0.1mol/L HClO ₄ 1mL=11.91mg C ₄ H ₉ NO ₃]
pН	5.2 to 6.2	AJI TEST 33
		[0.20g in 20mL of H ₂ 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]

 $^{^{1}\,\,}$ This product, in terms of actual quality, conforms to USP, EP, and JP.

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² Temperature coefficient of $[\alpha]_D^t$: +0.04°

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.

⁴ Disregard limit: 0.05%