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L-Citrulline-DL-Malate2:1 Specification Sheet

Item	Specification	Test Method
Description	White powder	Organoleptic
Identification	In accordance with standard	HPLC
Optical rotation (°)	+16.5~+18.5	ChP2015
Solubility	Clear	ChP2015
Loss on drying (%)	Not more than 0.3	ChP2015
Residue on ignition (%)	Not more than 0.1	ChP2015
Chloride (%)	Not more than 0.02	ChP2015
Sulfate (%)	Not more than 0.02	ChP2015
Ammonium (%)	Not more than 0.02	ChP2015
Heavy Metal (ppm)	Not more than 10	ChP2015
Lead (ppm)	Not more than 1	GB 5009.12-2017 I
Mercury (ppm)	Not more than 0.03	GB 5009.17-2014 I
Cadmium (ppm)	Not more than 0.1	GB 5009.15-2014
Arsenic (ppm)	Not more than 1	ChP2015
Iron (ppm)	Not more than 10	ChP2015
Assay (%)	Not less than 98.5	HPLC
L-Citrulline (%)	62.5~74.2	HPLC
DL-Malate (%)	25.8~37.5	HPLC
Bulk density (g/ml)	-----	ChP2015
Tightness (g/ml)	-----	ChP2015
30 Mesh (%)	-----	ChP2015
Total plate counts (cfu/g)	Not more than 1000	ChP2015
Yeast and Mould (cfu/g)	Not more than 100	ChP2015
Coliforms	Absence/g	ChP2015
E.coli	Absence/g	ChP2015
Salmonella	Absence/10g	ChP2015
Staphylococcus aureus	Absence/g	ChP2015

Conclusion: Qualified
 Analysis date: 2019-11-30 Analyst: _____ Verifier: 曹松影