



魯維制藥集團有限公司

Luwei Pharmaceutical Group Co.,Ltd.
Shuangfeng Industrial Park Zichuan District Zibo 255086 China
Tel: 0086 533 6220986 Fax: 0086 533 6220993

CERTIFICATE OF ANALYSIS

LWPC-1C1-21-07-069		Date: 15 OCT ,2021	
Product Name		ASCORBIC ACID	
Analysis Standard		BP/USP/EP/E300	
Batch No.		1210133097	
Quantity		22000kgs	
Manufacture Date		SEP 2021	
Expiry Date		AUG 2024	
Net/Gross Weight		25.0kgs/26.3kgs	
Analysis Contents		Analysis Standard	Analysis Results
Characteristics		White or almost White crystals Crystalline Powder	Pass
Identification		Positive Reaction	Positive
Melting Point		About 190°C	191.2°C
PH (5% aqueous solution)		2.1-2.6	2.35
PH (2 % aqueous solution)		2.4-2.8	2.54
Clarity Of Solution		Clear	Clear
Colour Of Solution		≤BY ₇	<BY ₇
Copper		≤5ppm	<5ppm
Heavy Metals		≤10ppm	<10ppm
Mercury		<0.1mg/kg	<0.1mg/kg
Lead		<2ppm	<2ppm
Arsenic		≤3ppm	<3ppm
Cadmium(Cd)		<1mg/kg	<1mg/kg
Oxalic Acid		≤0.2%	<0.2%
Iron		≤2ppm	<2ppm
Loss of Drying		≤0.4%	<0.4%
Sulphate Ash(Residue On Ignition)		≤0.1%	<0.1%
Specific Optical Rotation		+20.5° - +21.5°	+21.11°
Organic Volatile Impurities		Pass	Pass
Impurity E		≤0.2%	<0.2%
Related Substances	Impurity C	<0.15%	<0.15%
	Impurity D	<0.15%	<0.15%
	Unspecified	<0.1%	<0.1%
	Total	<0.2%	<0.2%



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Total Plate Count	100 cfu/g max	
E-coli	Negative	
Yeast and Mold	100 cfu/g max	
Pseudomonas	Negative	
Staphylococcus	Negative	
Salmonella	Negative	
Colour reaction	Pass test	
Reducing reaction	Pass test	
Assay	99.0%-100.5%	99.76%
Conclusion	The Above-Mentioned Product Conforms To BP/USP/EP/E300	

Approved by:Fengchunxiao

Rechecker:Jiajinfeng

Tester:Liping

For and on behalf of
鲁维制药集团有限公司
LUWEI PHARMACEUTICAL GROUP CO., LTD

Authorized Signiture(S)