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CERTIFICATE OF ANALYSIS

Product Name:	**Levodopa**		
Batch No.	C41-20230436	**Specifications**	EP10
Manufacturing Date	April 23, 2023	**Expiry Date TO**	April 22, 2026
Quantities/Spec.	502.83kg/25kg/drum	**Report Date**	May 12, 2023
Test Basis	SOP-FP-070-02	**Report No.**	C41-2023006(EP)
Analytical results			
Items	**Specifications**		**Results**
Appearance	White or almost white,crystalline powder		Almost white, crystalline powder
Solubility	Meets the requirements.		Meets the requirements.
Identification	A.IR The sample comparing with the spectrum obtained with standard		Complies
Appearance of solution	Clear and not more intensely coloured than reference solution BY6		Clear and less than BY6
pH	4.5~7.0		4.8
Related Substance	Impurity A:Not more than 0.1%		Not detected
	Impurity B:Not more than 0.5%		0.02%
	Impurity C:Not more than 0.2%		Not detected
	Unspecified impurities:Not more than 0.05%		0.01%
	Total impurities:Not more than 1.0%		0.05%
Enantiomeric purity	Not more than 0.5%		Not detected
Loss on drying	Not more than 1.0%		0.20%
Sulphated ash	Not more than 0.1%		0.02%
Assay	99.0%~101.0%		99.60%
Conclusion	The batch complies with the EP10 Specifications.		

Inspector: Carlifor Wu Verifier: Jack He

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