



东北制药集团股份有限公司
NORTHEAST PHARMACEUTICAL GROUP CO., LTD

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CERTIFICATE OF ANALYSIS

Sucralfate EP

BATCH NUMBER	DY02012500006	MANUFACTURE DATE	Mar.18.2025
BATCH SIZE	25kg	TEST DATE OF APPLICATION	Mar.25.2025
QUANTITY	1Drums	RETEST DATE	Mar.17.2028
Analysis Items		Specifications	Analysis Results
1.	Characteristics	White or almost white, amorphous powder	white, amorphous powder
2.	Identification	A:IR:Complies B:Positive C:Positive	Complies Positive Positive
3.	Acid Neutralization Equivalent	Meets the Requirements	Meets the Requirements
4.	Assay: (C ₁₂ H ₁₄ O ₁₀ S ₈)	30.0%~38.0%	33.9%
5.	Assay:Aluminum	15.5%~18.5%	17.9%
6.	Related Substances	Impurity A≤5.0%	0.04%
7.	Chloride	≤0.50%	<0.50%
8.	Particle-Size	100% pass though 50μm	100%

We, Northeast Pharmaceutical Group Co., Ltd., certify that this batch of **Sucralfate** meets the requirements of EP11.0

Supervisor

Final Batch Disposition

路阳 2025.04.01

Approved



By: 马红芳 2025.04.01

METAPHARMACEUTICAL

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13/05/2025