



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

Lot: 0020912
ID 221316

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CERTIFICATE OF ANALYSIS
Chloramphenicol USP/BP/EP

BATCH NUMBER	DY 00911242	MANUFACTURE DATE	Nov.26.2011
BATCH SIZE	1000 kg	TEST DATE	Nov.27.2011
QUANTITY	40 Drums	RETEST DATE	Nov.25.2016

Analysis Items		Specifications	Analysis Results
1.	Characteristics	Fine, white to greyish white or yellowish white, needle-like crystals or elongated plates	Yellowish white, needle-like crystals, elongated plates
2.	Identification	IR: Meets the requirements	Meets the requirements
3.	Melting Point	149°C ~153°C	149.9°C ~151.5°C
4.	Assay	98.5%~102.0%	99.2%
5.	Loss on Drying	≤0.5%	0.03%
6.	Residue on Ignition	≤0.1%	0.01%
7.	Acidity or Alkalinity (pH)	5.0~7.5	5.70
8.	Specific Rotation	+18.5°~+20.0°	+19.8°
9.	Related Substances	The any secondary spot≤0.5% Total≤2%	The any secondary spot<0.5% Total<2%
10.	Chloride	≤0.0050%	<0.0050%
11.	Crystallinity	Meets the requirements	Meets the requirements

We, Northeast Pharmaceutical Group Co., Ltd., certify that this batch of Chloramphenicol meets the requirements of United States Pharmacopoeia 34, British Pharmacopoeia 2011 and European Pharmacopoeia 7th.

Analysts 33180180

Checker

Supervisor

Final Batch Disposition

Approved

By:

