开封明仁药业有限公司

Add.:Kaifeng Mingren Pharmaceutical Co.,Ltd,

Bianjing Road East of embankment, Kaifeng City, Henan Province.475000

Tel: 86-371-65741762 Fax: 86-371-65706158

CERTIFICATE OF ANALYSIS

检验报告单

REGISTER No. 记录编码: RP-S-YC05001-00 REPORT No.:报告单编号: YC05001202210005

Product: Phloroglucinol Anhydrous	Batch No.:SB221001
产品名称: 无水间苯三酚	批号:
Manufacturing date:Sep.29,2022	Quantity:500.00kg
生产日期:	数量:
Retest Date:Aug.2025	Package:25kg/DRUM
复验期:	包装规格:
Sample Receiving Date:Oct.03,2022	Reporting Date:Oct.07,2022
收检日期:	报告日期:
Testing Item: Full test	Specification : Ph. Eur.10.0
检验项目: 全检	检验依据:

		157 457 IN AM.		
Items 检测项目 Method 方法		Acceptance Criteria 标准规定	Results 检验结果	
Appearance	Visual	White or almost white powder.	White powder	
性状	目测	本品为白色或类白色粉末。	白色粉末	
Solubility 溶解度	Visual 目測	Sparingly soluble in water, freely soluble in ethanol(96%),practically insoluble in methylene chloride 本品在水中略溶,在乙醇(96%)中易溶,在二氯甲烷中几乎不溶。	Conforms 符合规定	
pH pH 值	EP2.2.3	4.0 to 6.0 应为 4.0~6.0。	5.6	
Appearance of solution 溶液的澄清度与颜色	EP2.2.1 EP2.2.2	Clear and not more intensely coloured than reference solution BY5 溶液应澄清无色;如显色,与BY5 标准比色液比较,不得更深。	Conforms 符合规定	
Identification 鉴别 (a)IR 红外 (b)TLC 薄层色谱	EP2.2.24 EP2.2.27	(a)The infrared spectrum of the product should beconsistent with that of the control product anhydrous phloroglucinol CRS. 本品的红外光吸收图谱应与无水间苯三酚对照品的图谱一致。 (b)The principal spot in the chromatogram obtained with the test solution is similar in	Conforms 符合规定 Conforms 符合规定	
(c) Loss on drying 干燥失重	EP2.2.32	position and size to the reference solution 供试品溶液所显主斑点的位置和大小应与对 照品溶液的主斑点一致 (c) Maximum 1.0per cent 不得过 1.0%	0.2%	

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收检日期:		报告日期:
Testing Item: Full test		Specification : Ph. Eur.10.0
检验项目: 全检		检验依据:

Items 检测项目	Method 方法	Acceptance Criteria 标准规定	Results 检验结果
Related substances 有关物质	EP2. 2. 29	Impurity A not more than 1.5 times the principal peak in the chromatogram obtained with reference solution (0.15 per cent) 杂质 A 不得大于对照溶液色谱图中主峰的峰面积的 1.5 倍(0.15%)	Not detected 未检出
		Impurity D not more than 1.5 times the principal peak in the chromatogram obtained with reference solution (0.15 per cent) 杂质 D 不得大于对照溶液色谱图中主峰的峰面积的 1.5 倍(0.15%)	Not detected 未检出
		Impurity E not more than 1.5 times the principal peak in the chromatogram obtained with reference solution (0.15 per cent) 杂质 E 不得大于对照溶液色谱图中主峰的峰面积的 1.5 倍(0.15%)	Not detected 未检出
		Impurity K not more than 1.5 times the principal peak in the chromatogram obtained with reference solution (0.15 per cent) 杂质 K 不得大于对照溶液色谱图中主峰的峰面积的 1.5 倍(0.15%)	Not detected 未检出

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Testing Item: Full test			Specification: Ph. Eur.10.0	
检验项目: 全检			检验依据:	
Items 检测项目	Method 方法	Accepta	nce Criteria 标准规定	Results 检验结果
		principal with refe 杂质 L 面积的	L not more than 1.5 times the l peak in the chromatogram obtained erence solution (0.15 per cent) 不得大于对照溶液色谱图中主峰的峰 1.5 倍(0.15%)	Not detected 未检出
Related substances	EP2.2.29	peak in reference 杂质 I	y I not more than 1.5 times the principal the chromatogram obtained with e solution (0.15 per cent) 不得大于对照溶液色谱图中主峰的峰 1.5 倍(0.15%)	Not detected 未检出
有关物质		impurity chromato (0.1 per 对于任-	fied impurities impurity for each not more than the principal peak in the ogram obtained with reference solution cent) 一杂质,其峰面积应不得大于对照溶图中主峰的峰面积(0.10%)	Not detected 未检出
		in the ch solution 总杂质 ⁷	t more than 3 times the principal peak fromatogram obtained with reference (0.3 per cent) 不得大于对照溶液色谱图中主峰的峰 3 倍(0.3%)	Not detected 未检出

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检验项目: 全检			检验依据:		
Items 检测项目	Method 方法	Accepta	nce Criteria 标准规定	Results 检验结果	
Loss on drying	EP2.2.32	Maximu	m 1.0per cent	0.2%	
干燥失重		不得过 1.0%			
Chlorides	EP2.4.4	Maximum 200ppm		Conforms	
氯化物		不得过2	200ppm	符合规定	
G 1C +		3.7	500-	G C	

Loss on drying	EP2.2.32	Maximum 1.0per cent	0.2%
干燥失重		不得过 1.0%	
Chlorides	EP2.4.4	Maximum 200ppm	Conforms
氯化物		不得过 200ppm	符合规定
Sulfates	EP2.4.13	Maximum 500ppm	Conforms
硫酸盐		不得过 500ppm	符合规定
Sulfated ash	EP2.4.14	Maximum 0.1 per cent	0.05%
硫酸盐灰分		不得过 0.1%	0.03%
Residual solvent	GC	1,2,4-trimethyl benzene not more than 20ppm	Not detected
残留溶剂		1, 2, 4-三甲苯应不得过 20ppm	未检出
Content	EP2.2.20	99.0 per cent to 101.0 per cent (dried	
含量	101.	substance)	99.8%
		按干燥品计算,含间苯三酚(C ₆ H ₆ O ₃)为	99.070
	-	99 0%-101 0% 17 44 117 (- ++ 11. +-	777 // -7

Conclusion: The results conform with Ph. Eur.10.0

结论: 本批号产品符合 EP10.0

KAIFENG NINGREN PHARMACEUTICAL CO.,LTD

MEI Quality Manager: LIANG JUAN

Analyst: HAN JIAYUE

Reviewer: ZHU JUNMEI 复核人:

质量经理:

化验员: 复核