山东久隆信和药业有限公司

SHANDONG JIULONG SINHERO PHARMACEUTICAL CO.,LTD

分析报告单 Certificate of Analysis

品名	上酸头孢噻呋 上酸头孢噻呋	检验日期				
Product Name	Ceftiofur Hydrochloride	Analysis Date		JUL.22, 2024		
数 量		生产日期				
数 里 Batch Quantity	300kg	MFG. Date		JUL.18, 2024		
批号	01424071804	有效期	JUL		.17, 2026	
Batch No.		EXP. Date			~ = D (±) \	
分析项目	标准		分析结果		项目结论 Items	
Analysis Item	Standard	,		st results	Conclusion	
			类白色结晶性粉			
外 观	白色或类白色结晶性粉末			末	符合规定	
Appearance	Appearance White or off-white crystalline powder		Off-white crystalline powder		Conforms	
	供试品溶液主峰的保留时间应与对照品溶液主峰的保		orystanine powder			
鉴 别 Identification	留时间一致		符合规定		符合规定	
	The retention time for the major peak of sample should			onforms	Conforms	
	correspond to that of reference standard					
	本品的红外光吸收图谱应与对照品的红外图谱一致		符合规定 Conforms		符合规定	
	The infrared absorption spectrum of sample corresponds to that of reference standard				Conforms	
	甲醇溶液应显氯化物的鉴别反应		符合规定 Conforms		//× 人·柯户	
	Methanol solution should show positive reaction of				符合规定 Conforms	
11.24-12	chloride					
比旋度 Specific Optical Rotation	-115°∼-127°			-122°	符合规定 Conforms	
				2.6	符合规定	
PH	2.0-4.0				Conforms	
水 分	≤5.0%			3.2%	符合规定	
Water	25.070			3.270	Conforms	
炽灼残渣 Residue on Ignition	≤0.5%			0.1%	符合规定 Conforms	
重金属					符合规定	
Heavy Metals	≤20ppm		<20ppm		Conforms	
粒度分布	D90≤10 µ m		符合规定 Conforms		符合规定	
Particle Size	D70 < 10 F III				Conforms	
细菌内毒素 Bacterial Endotoxins	≤1.0EU/mg		<1.0EU/mg 符合规定		符合规定	
无 菌	符合规定				Conforms 符合规定	
Sterility	17 日 次及 Conforms		TT 市 规定 Conforms		Conforms	
有关物质(HPLC)	头孢噻呋肟乙酯应不得	过 0.5%	ND		符合规定	
Related Substances	Ceftiofur oxime ethyI ester should n	ot more than 0.5%			Conforms	

Address: North of Kaiyuan Street and West of Tianhe Road, Shanghe County Economic Development Zone, Jinan City, Shandong Province, China

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有关物质(HPLC) Related substances	7-氨基头孢烷酸应不得过 0.5% 7-Aminocephalosporanic acid should not more than 0.5%	ND	符合规定 Conforms		
	2-呋喃甲酸应不得过 0.5% 2-Furoic acid should not more than 0.5%	ND	符合规定 Conforms		
	N-去乙酰头孢噻呋应不得过 0.5% N-Deacyl ceftiofur should not mnore than 0.5%	ND	符合规定 Conforms		
	头孢噻肟应不得过 0.5% Cefotaxime should not more than 0.5%	0.01%	符合规定 Conforms		
	头孢噻呋 δ -3 异构体应不得过 0.5% Ceftiofur delta-3 isomer should not more than 0.5%	0.01%	符合规定 Conforms		
	头孢噻呋 E-异构体应不得过 3.6% Ceftiofur E-isomer should not more than 3.6%	0.03%	符合规定 Conforms		
	双氢噻吩硫酯应不得过 0.5% Dihydrothiophenyl thioester should not more than 0.5%	0.01%	符合规定 Conforms		
	头孢噻呋氨基二聚物应不得过 0.8% Ceftiofur amide dimer should not more than 0.8%	ND	符合规定 Conforms		
	N-三苯甲基头孢噻呋肟应不得过 0.5% N-Trityl ceftiofur oxime should not more than 0.5%	ND	符合规定 Conforms		
	其他单一杂质应不得过 0.5% Other single impurities should not more than 0.5%	符合规定 Conforms	符合规定 Conforms		
	杂质总和应不得过 6.0% Total of the impurities should not more than 6.0%	0.9%	符合规定 Conforms		
含 量 Assay(HPLC) (以无水物计 calculated on the anhydrous basis)	含头孢噻呋不得少于 85.0% Contain Ceftiofur should not less than 85.0%	92.2%	符合规定 Conforms		
	含盐酸头孢噻呋不得少于 91.0%。 Contain Ceftiofur HCl should not less than 91.0%	98.6%	符合规定 Conforms		
结论 Conclusion	本品符合 CVP2020。 Conclusion: Complies with the requirements of CVP2020.				

Analyst:

质保部批 Approved by