

CONCORD BIOTECH LIMITED

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CERTIFICATE OF ANALYSIS

Product		CICLOSPORIN Ph.Eur.				L- 13418095B	
Batch No.		13418095		Mfg. Date	Nov	Nov 2018	
Batch size		30.240 Kg	0.240 Kg		Oct	2022	
S.N.	Tests Specifications Observations						
1			White or all	White or almost white powder.		White powder,	
2	Solubility		Anhydrous	Practically insoluble in water, freely soluble in Anhydrous ethanol and in Methylene chloride.		Practically insoluble in water, freely soluble in Anhydrous ethanol and in Methylene chloride.	
3	Identification A. [By IR] B. [By HPLC]		The absorption maxima in the spectrum obtained with the substance to be examined corresponds in position and relative size to those in the spectrum obtained with Ciclosporin reference standard / working standard.			The absorption maxima in the spectrum obtained with the substance corresponds in position and relative size to those in the spectrum obtained with Ciclosporin working standard.	
			The principal peak in the chromatogram obtained with the test solution is similar in retention time to the principal peak in the chromatogram obtained with reference solution (a) prepared under the assay test.			Std RT: 27.3 minutes Test RT: 26.9 minutes	
4	Appearance of solution		The solution is clear and not more intensely colored			The solution is clear.	
•			than referen	ce solution Y ₅ , BY ₅ or R ₇ .	Complies		
5	Specific optical rotation		Between -193° and -185°		-187°		
	(Calculated with reference to the dried substance)						
6	Loss on Drying		Not more than 2.0 %		0.53 %		
7	7 Related substances [By HPLC]						
	(a) Ciclos		Not more th	1700TO (B) B (C) 1700B		Below disregard limit Below disregard limit	
	(b) Ciclosporin B		Not more than 0.7 %		Below disregard limit Below disregard limit		
	(c) Ciclosporin L		Not more than 0.7 % Not more than 0.7 %		Below disregard limit		
	(d) Ciclosporin U (e) Ciclosporin H		Not more than 0.7 %		Not detected		
		rociclosporin A		Not more than 0.7 %		0.29%	
				Not more than 0.7 %		0.11%	
	(g) Ciclos			Not more than 0.7 %		0.16%	
	(h) Ciclos (i) Ciclos	porin E	Not more th			Not detected	
	(i) Inopini	losporin A	Not more th			Below disregard limit	
	(b) Apy in	dividual unknown impurity	Not more th			Below disregard limit	
		f all impurities	Not more th			0.56%	
8	Assay [By			2.0% (as C ₆₂ H ₁₁₁ N ₁₁ O ₁₂).		100.2 %	
o	(on the dri			your Theoretic of Tiley.		in the state of th	
Addition	Additional Tests:						
9	Residual solvents (By GC HEADSPACE)						
,	Acetone		NMT 4500	NMT 4500 ppm		11 ppm	
	Ethyl acet	ate	NMT 2000			Below detection limit	
10	Particle size distribution			For information only		d(0.9): 55 microns	
11	Microbial EnumerationTests:						
	Total viab	le aerobic count		NMT 1000 CFU / g		Less than 10 cfu / g	
	Total mole	d and yeast count .	NMT 100 C	NMT 100 CFU / g		Less than 10 cfu / g	
	Test for specified microorganism			10 No. 10			
	E. coli			Should be Absent / g		Absent / g	
	S. aureus		Should be A			Absent / g	
	Pseudomo	nas aeruginosa '	Should be A			Absent / g	
	Candida a		Should be A			Absent / g	
		m sporogenes	Should be A			Absent / g	
	Salmonella spp.		Should be A	Should be Absent /10 g		Absent / 10g	

Storage: Store in an airtight container, protected from light upto 25°C.

Remarks: The material complies as per Ph. Eur and in-house specification no. SPC/QC/FP/033-03.

Date of Report: Dec 04, 2018

Compiled By (QC)

(D.N.Panchal-Sr.AM)

Date:

Checked By (QC)
(N.S.Jagad-DGM)
Date:

Approved By (QA) (T.K.Saha-DGM