台州天禄生物医药有限公司

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CERTIFICATE **ANALYSIS** OF

Name of product	Eprinomectin	MFG. Date	April.28,2018
Batch No.	TYL20180428	EXP. Date	April.27,2020
QTY.	52kg	Test Standard	USP
Test Items	Specifications		Result
Appearance	white or yellowish-white crystalline powder		Conforms
Identification	IR:Corresponds with that of standard		Conforms
	HPLC:The retention time for B _{1a} and B _{1b} Corresponds with that of standard		Conforms
Water	≤2.0%		0.70%
Limit of 8a-oxo-B1a	≤0.5%		0.40%
Heavy Metals	≤10ppm		Conforms
Residue on ignition	≤0.1%		0.03%
Acetonitrile	≤0.005%		0.002%
The sum of all solvents	≤0.5%		0.42%
Related substances	Relative retentions of 0.23,0.93,1.16≤1.0%		0.02%,0.02%,0.02%
	Impurity A≤1.0%		0.23%
	Impurity E≤1.0%		0.03%
	Any other known Impurity≤0.5%		0.216%
	Total unknown Impurities≤1.0%		0.18%
	Total Impurity≤5.0%		0.76%
Assay(on dried basis)	B₁≥95.0%		98.65%
	B _{1a} ≥90.0%		96.13%
Conclusion	It complies with all requirements of USP standard		

QC.Manager:叶晓培

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Analysed by:于明明

