

ALIVIRA ANIMAL HEALTH LIMITED CERTIFICATE OF ANALYSIS



PONAZURIL

Batch No.	ASA002726A	Quantity	150.00 Kg
Manufacturing Date	August-2021	Retest/-Expiry Date	July-2024
A.R.NO.	ALIV-FP-210493	Lot No.	
Storage Condition	Store in tight container at 25°C, excursion permitted between 15 to 30°C		
Customer Name/ Vender Name	SUAN PHARMA		

S.No.	TEST PARAMETERS	SPECIFICATIONS	DECEMBER OF
1	Description	White to almost white powder.	RESULTS
2	Identification by IR	The infrared absorption spectrum is concordant with spectrum obtained from standard	White powder Complies
3	Identification by HPLC	The retention time of sample corresponds to that of retention time of standard as obtained in assay by HPLC	Complies
4	Solubility	Sparingly soluble in Ethyl acetate	Comple
5	Apperance of solution	The solution is clear and not more intensely colored than B7 and Y7	Complies Complies
6	Loss on Drying	Not more than 1.00% w/w	0.24%
7	Water content	Not more than 0.50% w/w	0.34%
8	Residue on ignition	Not more than 0.10% w/w	0.05%
9	Heavy metals	Not more than 10 ppm	Complies
10	Related substance by HPLC Toltrazuril Sulfoxide N-Methyl Toltrazuril Sulfoxide N-Methyl Toltrazuril sulfone Toltrazuril Any individual impurity Total impurities	Not more than 0.50% Not more than 0.20% Not more than 0.20% Not more than 0.50% Not more than 0.20%	0.05% Not detected 0.03% Not detected 0.08% 0.25%

	PREPARED BY	REVIEWED BY	APPROVED BY
DESIGNATION	Sr.Executive-QC	Head-QC/Designee	
SIGNATURE	LN	(4)	SA 2
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Format No.: QC/116/FMT-001/R0

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Corporate Office: 301, Dosti Pinnacle, Plot No.22, Wagle Indl. Area, Thane (W)-400 604, India. T+912241114777 Factory Address: Plot No.104 to 109, Part of 112 & 113, Ramky Pharma City India Limited -SEZ, JNPC, Parawada Mandal, Visakhapatnam – 531 019, Andhra Pradesh, India. Email: info@alivira.in Web: www.alivira.co CIN: U74120MH2013PLC248708



ALIVIRA ANIMAL HEALTH LIMITED CERTIFICATE OF ANALYSIS



PONAZURIL

Batch No.		ASA002726A	Quantity	150.00 Kg	
Manufacturing Date		August-2021	Retest/Expiry Date	July-2024	
A.R.NO.		ALIV-FP-210493	Lot No.	40000052734	
	e Condition		at 25°C, excursion permi	tted between 15 to 30°C.	
Customer Name/Vendor Name		SUAN PHARMA			
S.No. TEST PARAMETERS		SPECIFICATIONS		RESULTS	
11	Assay by HPLC (On dried basis)	Between 98.0% to 102.0%		99.4%	
12	Acetic Acid Content by HPLC	Not more than 1000 ppm		Not detected	
13	Residual solvents by GC Dimethyl sulphoxide Methanol Benzene	Not more than 5000 ppm Not more than 3000 ppm Not more than 2 ppm		Not detected 27 ppm Not detected	
Rema	rks : The above batch complies a	ns per IH Specifications	el minima pilli.	anticipals to surrough)	
		PREPARED BY	REVIEWED BY	APPROVED BY	
DESIGNATION		Sr.Executive-QC	Head-QC/Designee	Head-QA/Designec	
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DATE	3	02/08/20	n 3238 2011	- Coxposis	
	at No.: QC/116/FMT-001/R0			Page 2 of 2	

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