



CENTURION REMEDIES PVT. LTD.

G-5, INDUSTRIAL ESTATE, GORWA, VADODARA-390016
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A/C OF HEALING PHARMA INDIA PVT LTD

QUALITY CONTROL DEPARTMENT

THE DRUG & COSMETIC ACT. 1940 & THE RULES THEREUNDER

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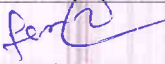
FINISHED PRODUCT CERTIFICATE OF ANALYSIS

Product Name : IVERHEAL-3	A.R. No. : F/2145/21-22
Packing : 10x10 TAB	Rel. Dt. : 02-09-2021
Generic Name : IVERMECTIN TABLETS USP 3 MG	T.R. Slip No. : CHPAP10109
Product Code : TI-53	T.R. Slip Dt. : 02-09-2021
Batch No. : TI-530003	Analysis Date : 02-09-2021
Actual Batch Size : 303000 TAB	Specification No. : CRFS/TI-53/21-22
Sample Size : 50.000 TAB	Specification Dt. : 13-04-2021
Released Qty : 303000.000 TAB (3030 CAR)	STP No. : STP/TI-53/AS PER USP
Remarks : Pass	Location : GORWA01
Mfg. Dt. : 08/2021	Make : HEALING
Exp. Dt. : 07/2024	
Test Packing : 10X10	
Mfg. Lic No. : G/25A/5141-A	
Test As Per : USP	

Sr.	Test	Result	Specification
1	DESCRIPTION	A white coloured, round shaped, Standard Concave, uncoated tablet ,plain on both side.	A white coloured, round shaped, Standard Concave, uncoated tablet ,plain on both side.
2	IDENTIFICATION	Complies.	Must Complies as per USP
1	BY HPLC	in the chromatogram of the Assay preparation correspond to those in the chromatogram of the Standard preparation, as obtained in the Assay	in the chromatogram of the Assay preparation correspond to those in the chromatogram of the Standard preparation, as obtained in the Assay
3	AVERAGE WEIGHT OF TABLETS	98.1 mg	97.00 mg \pm 5.0% of avg.weight.
4	UNIFORMITY OF WEIGHT	H.L. : + 2.73 % L.L. : - 2.56 %	Within \pm 7.5% of avg.weight.
5	DIAMETER	6.03 mm	6.00 mm \pm 0.2 mm
6	THICKNESS	3.04 mm	3.00 mm \pm 0.3 mm
7	HARDNESS	4 Kg/cm ²	NLT 3 Kg/cm ²
8	FRIABILITY	0.41 %	NMT 1.0 %
9	DISINTEGRATION TIME	06 Minute 53 Seconds in water Medium at 37°C	NMT 15 minutes in water medium at 37°C
10	DISSOLUTION	100.01%	NLT 80.0 % (Q)
11	UNIFORMITY OF DOSAGE UNITS	Min : 102.41 % Max : 108.95% Avg : 105.03%	85.0 % TO 115.0 %
12	ASSAY	100.86%	Each uncoated tablets contains ; Ivermectin IP 3 mg [Limit ; 90.0% To 110.0%]

Conclusion : The above sample complies as per USP

In the Opinion of the undersigned the sample referred to above is of Standard quality as defined in the Act and the Rules made thereunder for the result given above. "This computer generated certificate of analysis is valid without signature"

Prepared By  PARTHSAPARA QC CHEMIST	Checked By PINALPATEL Q.C IN CHARGE	Approved By PUSHPENDRA QA HEAD
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