



Saintlife Pharmaceutical Ltd.

(An ISO -9001-2015/QMS Cert.Co.) GMP & WHO Certified Company.
323, Central hope -Town, industrial Area, Selaqui, Dehradun -248011 (U.K.)

CERTIFICATE OF FINISHED PRODUCT OF ANALYSIS

(The Drugs & Cosmetics Act.1940 & Rules there under)

Manuf.Lic.No.34/UA/2017

(Form -39)

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33/UA/SC/P-2017

Name of Product	IVERJOHN-3	A.R. No.	FP-100/03-2021
Generic Name	IVERMECTIN TABLETS USP	Sample Quantity	60 Tablets
Batch No.	TA21053	Batch Size	10.0 Lacs
Date of Mfg.	03/2021	Date of Exp.	02/2024
Date of Receipt	15/03/2021	Date of Release	16/03/2021

S.No.	Test Parameter	Specification	Results
1.	Description	White colour, round shape, biconvex, having break line on one side, plain on other side and uncoated tablets. 10X10 tablets packed in a carton.	White colour, round shape, biconvex, having break line on one side, plain on other side and uncoated tablets. 10X10 tablets packed in a carton.
2.	Identification	Should be positive for ivermectin as per assay procedure.	Positive for ivermectin as per assay procedure
3.	Average weight	135.0 mg \pm 5.0 %	134.16 mg
4.	Uniformity of weight	135.0 mg \pm 7.5 %	-1.7 % to +2.5%
5.	Disintegration Time	Not more than 15 minutes	1.15 minutes
6.	Hardness	Not less than 2.0 Kg	2.90 Kg
7.	Thickness	3.20 mm \pm 0.20 mm	3.28 mm
8.	Diameter	6.50 mm \pm 0.20 mm	6.49 mm
9.	Friability	Not more than 1.0 %	0.14 %
10.	Dissolution	Not less than 80 % (Q) of the labeled amount of the Ivermectin.	98.17% to 99.71%



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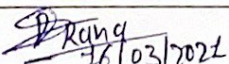

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11.	Assay: Each uncoated tablet contains: Ivermectin USP 3.0 mg	2.7 mg to 3.3 mg (90.0% to 110.0%w/w of Label Claim)	2.999 (99.97%)
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Opinion: The above material is complies / does not comply with the specification.

	Analysed By	Approved By
Sign/Date	 16/03/2021	 16/03/2021
Name	Poonam Rana	Sheetal Sharma
Designation	Analyst QC	Sr. Officer QC

