



## 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.)  
Mobile No.:9967006091, E-mail:admin@johnleeindia.com

### 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-12	A.R. No.	FP-0183/05-2021
Generic Name	IVERMECTIN TABLETS USP	Sample Quantity	60 Tablets
Batch No.	TA21104	Batch Size	1340000
Date of Mfg.	04/2021	Date of Exp.	03/2024
Date of Receipt	01/05/2021	Date of Release	03/05/2021

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	130.0 mg $\pm$ 5.0 %	131.07mg
4.	Uniformity of weight	130.0 mg $\pm$ 7.5 %	-2.57 % to +2.69%
5.	Disintegration Time	Not more than 15 minutes	30 seconds
6.	Hardness	Not less than 2.0 Kg	3.4 Kg
7.	Thickness	3.20 mm $\pm$ 0.20 mm	3.32 mm
8.	Diameter	6.50 mm $\pm$ 0.20 mm	6.61 mm
9.	Friability	Not more than 1.0 %	0.06 %
10.	Dissolution	Not less than 80 % (Q) of the labeled amount of the Ivermectin.	92.25% to 99.58%



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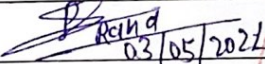
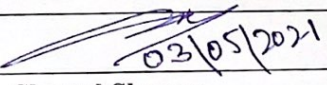
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11.	Assay: Each uncoated tablet contains: Ivermectin USP 12mg	10.80 mg to 13.20 mg (90.0% to 110.0%w/w of Label Claim)	11.984 mg (99.86 %)
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Opinion: The above material is complies / ~~does not comply~~ with the specification.

	Analysed By	Approved By
Sign/Date	 03/05/2021	 03/05/2021
Name	Poonam Rana	Sheetal Sharma
Designation	Analyst QC	Sr. Officer QC

