



20 APR 2022

FDA ADVISORY  
No. **2022-0942**

**TO : ALL HEALTHCARE PROFESSIONALS AND THE GENERAL PUBLIC**

**SUBJECT : Public Health Warning Against the Purchase and Use of the Unregistered Drug Product “Ivermectin Tablet USP Iverjohn-12 12 mg Tablet”**

The Food and Drug Administration (FDA) advises the public against the purchase and use of the unregistered drug product:

**Composition :**  
Each uncoated tablet contains :  
Ivermectin 12 mg  
Excipients q.s.  
**Dosage :** As directed by Physician.  
**Storage :** Store in a cool, dry & dark place.  
Keep the medicine out of reach of children.

Manufactured in India for  
**Johnlee Pharmaceuticals Pvt. Ltd.**  
Saintlife Pharmaceuticals Ltd.  
121, Central Hope-Town Industrial Area, Selaqui Dehradun-248011 (U.K.)  
Marketed By  
**Johnlee**  
Johnlee Pharmaceuticals Pvt. Ltd.  
42/44, Babu Genu Road,  
Om Shanti Co-Operative Hsg  
Society, 2nd Floor, Shop No-6,  
Kalbadevi, Mumbai-400 002  
Email : admin@johnleeindia.com

Mfg. Lic. No. : 34/UA/2017  
33/UA/SC/P/2017  
Batch No. : TA21168  
Mfg. Date : 08/2021  
Exp. Date : 07/2024  
M.R.P. : 350.00  
For 10 Tablets  
Incl. of all taxes

**IVERMECTIN TABLET USP**  
**IVERJOHN-12**  
10 x 10 Tablets  
12 mg

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Email : admin@johnleeindia.com

**“Ivermectin Tablet USP Iverjohn-12 12 mg Tablet”**

Manufactured by: Saintlife Pharmaceuticals Ltd. – 323, Central Hope-Town Industrial Area, Selaqui Dehradun-248011 (U.K.)

Marketed by: Johnlee Pharmaceuticals Pvt. Ltd. 42/44, Babu Genu Road, Om Shanti Co-Operative Hsg Society, 2<sup>nd</sup> Floor, Shop No-6, Kalbadevi, Mumbai-400 002

Figure 1: Unregistered drug product



FDA Post-Marketing Surveillance (PMS) activities have verified that the abovementioned drug product has not gone through the registration process of the Agency and has not been issued with proper authorization in the form of Certificate of Product Registration. Thus, the Agency cannot guarantee its quality and safety. Therefore, consumption of such violative product may pose potential danger or injury to health.


Pursuant to Republic Act No. 9711, otherwise known as the "Food and Drug Administration Act of 2009", the manufacture, importation, exportation, sale, offering for sale, distribution, transfer, promotion, advertising or sponsorship of health products without proper authorization from FDA is prohibited.

All concerned establishments and/or entities are warned not to distribute the above-identified violative drug product until it has been covered by the appropriate authorization, otherwise, regulatory actions and sanctions shall be strictly pursued. Always check if a product is registered with the FDA by using the **FDA Verification Portal feature** accessible at <https://verification.fda.gov.ph>.

All Local Government Units (LGUs) and Law Enforcement Agencies (LEAs) are requested to ensure that this product is not sold or made available in their localities or areas of jurisdiction.

For more information and inquiries, please e-mail us at [info@fda.gov.ph](mailto:info@fda.gov.ph). To report continuous sale or distribution of unregistered health products, kindly e-mail us via [ereport@fda.gov.ph](mailto:ereport@fda.gov.ph). You may also call the Center for Drug Regulation and Research at telephone number (02) 8809-5596. For any suspected adverse drug reaction (ADR), report immediately to FDA through this link: <https://primaryreporting.who-umc.org/Reporting/Reporter?OrganizationID=PH> and fill out all the required fields.

Dissemination of the information to all concerned is requested.

  
**DR. OSCAR G. GUTIERREZ, JR.**  
Officer-in-Charge Director General

DTN:



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