TruPharma™

January 18, 2022

MANUFACTURED BY:
Rubicon Research Private Limited
Ambernath, Dist. Thane, 421506 India

DISTRIBUTED BY:
TruPharma, LLC
Tampa, FL 33609

Dear Healthcare Partner.

URGENT: DRUG RECALL – RETAIL LEVEL

Metoprolol Tartrate Tablet USP, 25mg 1000

TruPharma, LLC (TruPharma) is initiating a voluntary recall of Lot **210211Hl** Expiry February 2024 of Metoprolol Tartrate Tablets USP, 25mg 1000 to the **retail level**. Rubicon Research Private Limited (Rubicon) manufactured and supplied the product for distribution by TruPharma. As an abundance of caution, this lot is being recalled due to a market complaint pertaining to foreign matter contamination of the lot.

A single tablet was identified to have a foreign object, and Rubicon has conducted an investigation into the source of the contamination and determined that this is an isolated, one-off incident.

Immediately examine your inventory and quarantine the product lot subject to the recall. Wholesalers and distributors should forward this notification to the retailers. Wholesalers and distributors who have the affected lot in inventory should contact Inmar Rx Solutions (Inmar) at 1-855-319-5713 Monday–Friday 9:00 a.m. to 5:00 p.m. EST. For reimbursement, please have the recalled lot returned to Inmar Rx Solutions on or before April 13, 2022. The lot number of the product can be found on the side of the bottle.

Metoprolol Tartrate Tabs USP, 25mg 1000

Strength	Lot #	NDC	Pack Size	Description
25mg	210211Hl	52817-360-00	1000	Metoprolol Tartrate 25mg tablets are film-coated, pink colored, round, biconvex tablets debossed with R 25 on one side and scored on the other side.

See enclosed Product Label for ease in identifying the product at **retail level**.

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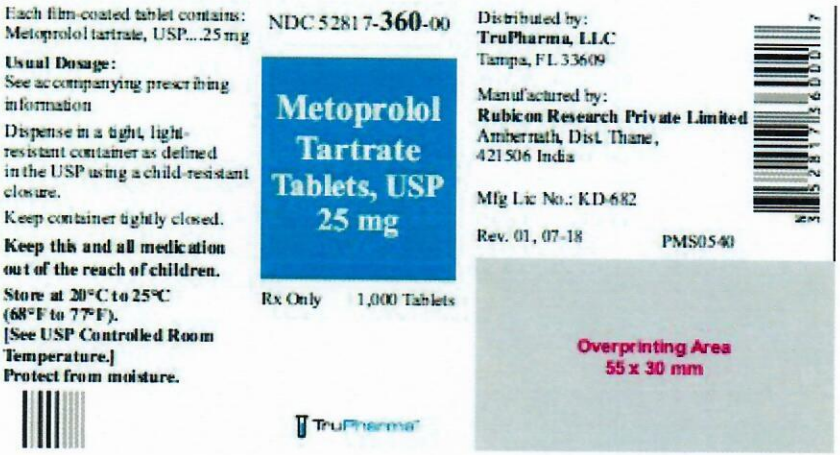

A COMPLETE PACKAGE OF INFORMATION INCLUDING A REPLY FORM WILL BE MAILED WITHIN (5) BUSINESS DAYS. THE REPLY FORMS SHOULD BE RETURNED TO INMAR THROUGH MAIL/EMAIL/FAX. ONCE RECEIVED A RETURN AUTHORIZATION AND BOX LABEL WILL BE PROVIDED

Upon receipt of this packet, please take the following actions:

1. Distributors/Pharmacies – Immediately examine your inventory, quarantine, and discontinue distribution of this lot.
2. Distributors – Complete the enclosed Business Recall Response Form even if you do not have any product on hand.
3. Distributors – Please pass this Recall Notice **ONLY** to pharmacies that received this product lot.
4. Pharmacies – If you have units of the affected products/lot in inventory, please contact Inmar at (855) 319-5713 to receive a Business Recall Response Form or acquire it from www.clsnetlink.com.
5. Business Recall Response Forms can be submitted by any of these methods.
   a. Fax: 817-868-5362
   b. Email: rxrecalls@inmar.com
   c. Mail: Inmar, Attn: Recall Coordinator - 635 Vine St, Winston Salem, NC 27101
6. Distributors/Pharmacies – Return recalled product to Inmar as instructed in recall/return packet.
7. Pharmacies – You do not need to contact any patients.

Upon receipt of the completed BRF, a return kit will be sent with an Return Authorization from and necessary box labels.

We appreciate your immediate attention to this matter. This recall is being made with the knowledge of the U.S. Food and Drug Administration.

Sincerely,

JB Davis
Chief Commercial Officer