



URGENT VOLUNTARY RECALL: Pharmacy Level, September 16, 2025

Levothyroxine Sodium Tablets USP 88 mcg

Accord Healthcare, Inc. (“Accord Healthcare”) is voluntarily recalling one lot of Levothyroxine Sodium Tablets USP 88 mcg, at the Pharmacy Level.

This recall is being initiated because out of specification results were observed during long term stability testing of Levothyroxine Sodium Tablets USP for 88 mcg. The out-of-specification results were observed on an assay conducted on one lot of Levothyroxine Sodium Tablets.

The assay content of Levothyroxine Sodium Tablets for lot D2300045 (88 mcg, 94.9% at 30 months) was observed below the approved specification range of 95.0%–105.0% set forth on the label. This means that the observed level of active ingredients in the product was below the approved specification.

These may affect the efficacy of the medication. As patient safety is the highest priority, Accord Healthcare is taking immediate action to recall the affected product lot.

Please examine your inventory of Accord Healthcare’s Levothyroxine Sodium Tablets USP 88 mcg for the below listed lot number carefully.

The product label for the recalled product should have the following details, please also refer to the enclosed product label included with this recall letter.

Levothyroxine Sodium Tablets USP				
Strength	Bottle Pack size	Product NDC	Lot No.	Expiry Date
88 mcg	90 Tablets	16729-450-15	D2300045	12/2025

Pharmacy - Please perform the following activities:

- Examine your inventory immediately for the listed lot number of Levothyroxine Sodium Tablets USP 88 mcg.
- Immediately discontinue distribution of the recalled Lot number of Accord Healthcare’s Levothyroxine Sodium Tablets USP 88 mcg.
- Promptly complete the attached Product Recall Response Form and reply even if you have **NO** Product to return.
- If you do have product to return, complete the attached Product Recall Response Form, quarantine the stock and follow the instructions given on the recall response form.
- If you have further distributed this lot number to other retailers, please immediately contact them and advise them of the recall and have them return their outstanding recalled stock to you. Return this stock as per the instructions on the attached Product Recall Response Form.





Your assistance is appreciated and necessary to prevent any potential health risk to the consumer.

Accord is working with Inmar Inc. to complete this recall. This recall is being made with the knowledge of the Food & Drug Administration. Your cooperation and prompt response to this notice is much appreciated.

Please complete and return the enclosed “PRODUCT RECALL RESPONSE FORM” as soon as possible, but no later than five business days from receipt of this letter.

**Completed Product Recall Response form should be emailed, or sent via FAX to INMAR,
Attn: Inmar Rx Solutions, 3845 Grand Lakes Way, Grand Prairie, TX 75050.
INMAR Email: rxrecalls@inmar.com. FAX: 1-817-868-5362.**

If you have any questions about the logistics for returning affected lot or other issues, please call Recall Services at 1-855-596-9369, Monday – Friday (excluding holidays), 9am to 5 pm EST.

INMAR will send you a Return Goods Authorization and shipping label. Appropriate credit for the returned product plus handling and shipping expenses will be issued to you upon receipt of the recalled product with the completed Return Goods Authorization. All recalled products returned without a Return Goods Authorization may delay the issuance of your credit.

We appreciate your assistance in this matter.

Sincerely,

A handwritten signature in blue ink, appearing to read "Sabita Nair", written over a light blue horizontal line.

Sabita Nair, RAC, ASQ-CPGP
Vice President – Regulatory Affairs
Accord Healthcare, Inc.
8041 Arco Corporate Drive, Suite 200
Raleigh, NC 27617, USA



Each tablet contains 88 mcg (0.088 mg) levothyroxine sodium USP.

See package insert for full prescribing information.

Contains FD&C Yellow No. 5 (tartrazine) as a color additive.

Store at 20°C to 25°C (68°F to 77°F); excursions permitted to 15°C to 30°C (59°F to 86°F) (see USP Controlled Room Temperature). Protect from light and moisture.

Dispense in a tight, light-resistant container as described in USP.

Do not accept if seal over bottle opening is broken or missing.

Manufactured for: Accord Healthcare, Inc.,
Raleigh, NC 27617.

Manufactured by: Intas Pharmaceuticals Limited,
Camp Road, Selaqui, Dehradun-248 197, INDIA.

NDC 16729-450-15

Levothyroxine Sodium Tablets, USP

88 mcg
(0.088 mg)

Rx Only

90 Tablets

accord

Mfg. Lic. No.: 15/UJ/SC/P-2006
80 3557 2 8618924 INL087



Keep area blank & varnish free for
overcoding Lot and EXP & Data matrix
40 X 20 mm



PRODUCT RECALL RESPONSE FORM

Product Recall Date: September 16, 2025

Voluntary Recall: Pharmacy Level

Levothyroxine Sodium Tablets USP				
Strength	Bottle Pack size	Product NDC	Lot No.	Expiry Date
88 mcg	90 Tablets	16729-450-15	D2300045	12/2025

Please fill out this form completely. By doing so, this will acknowledge that you have read and understand the recall instructions and have taken the appropriate action.

Customer Name _____ DEA # _____
**DEA # is required, if not provided the processing of your form will be delayed.*

Address _____

City _____ State _____

Zip _____

Contact Name (please print) _____ Telephone # _____

Contact Signature _____ Date _____

If you did not purchase the product directly from the Manufacturer, please complete the following section.

Purchased from: Name _____ DEA # _____

Address _____

City _____ State _____

Zip _____

Please check all appropriate boxes:

- I have read and understand the recall instructions provided in the letter.
- I have checked my stock and have quarantined inventory consisting of _____ bottles/units.

Any adverse events associated with recalled product?

Yes NO If yes, please explain: _____

Please describe your business: _____



I have checked my stock and:

_____ Do not have any stock of recalled **items**.

OR

_____ Have quarantined and listed in the box above the quantity of bottles/units of **Levothyroxine Sodium Tablets USP 88 mcg**, and will be returning them to Inmar, as soon as possible.

Upon receipt of this Response Form, Inmar will issue return authorization label(s).

Please indicate the number of box labels needed: _____

Completed Product Recall Response form should be emailed, or sent via FAX to INMAR,
Attn: Inmar Rx Solutions, 3845 Grand Lakes Way, Grand Prairie, TX 75050.
INMAR Email: rxrecalls@inmar.com. FAX: 1-817-868-5362.

Even if you do not possess any inventory of the lot being recalled, we would appreciate it if you could still fill out and return the “PRODUCT RECALL RESPONSE FORM”.

If you have any questions about the logistics for returning affected lot or other issues, please call Recall Services at 1-855-596-9369, Monday – Friday (excluding holidays), 9am to 5 pm EST.

