



# Lupin Pharmaceuticals, Inc.

## URGENT: DRUG RECALL

April 24, 2026

On behalf of Lupin Pharmaceuticals Inc.  
Inmar Rx Solutions, Inc.,  
Attn: Recall Coordinator,  
One West Fourth Street, Suite 500 Winston Salem, NC 27101.

Dear Healthcare Partner,

This is to inform you of a product recall involving:

### Liraglutide Injection 6 mg/mL (2s Pack)

Strength	Lot	Expiry	NDC	Description
6 mg/mL	WB00097	Jul. 2027	70748-346-02	Clear colorless solution free from visible particulate matter in glass cartridge assembled in blue colored pen injector device with blue button.
	WB00094	Jul. 2027		
	WB00088	Jul. 2027		
	WB00103	Oct. 2027		
	WC00016	Dec. 2027		

### Liraglutide Injection 6 mg/mL (3s Pack)

Strength	Lot	Expiry	NDC	Description
6 mg/mL	WB00098	Jul. 2027	70748-346-03	Clear colorless solution free from visible particulate matter in glass cartridge assembled in blue colored pen injector device with blue button.
	WB00092	Jul. 2027		
	WB00093	Jul. 2027		
	WB00104	Oct. 2027		
	WB00106	Oct. 2027		
	WB00105	Oct. 2027		

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[www.lupin.com/us](http://www.lupin.com/us)



# Lupin Pharmaceuticals, Inc.

As a precautionary measure, Lupin Pharmaceuticals, Inc. is initiating a **voluntary recall** of five lots of **Liraglutide Injection 6 mg/mL (2s Pack)** and six lots of **Liraglutide Injection 6 mg/mL (3s Pack)** at the **Retail** level. These drug product lots are being recalled due to a single market complaint. The reported complaint was identified as a white thread-like structure in the cartridge. Based on the Health Hazard Assessment conducted for the reported complaint, no adverse events are expected from the consumption of these products. In a worst-case scenario, the product may have the potential to cause acute, localized inflammatory reactions.

See enclosed product label(s) for ease of identifying the product.

The recalled lots were distributed nationwide to wholesalers, distributors, supermarkets and mail order pharmacies between October 2025 and April 2026.

**A COMPLETE PACKAGE OF INFORMATION INCLUDING A REPLY FORM WILL BE MAILED WITHIN (5) BUSINESS DAYS. THE REPLY FORMS SHOULD BE RETURNED TO INMAR Rx SOLUTIONS, INC. THROUGH MAIL/EMAIL/FAX. ONCE RECEIVED AN RA AND NEEDED BOX LABELS WILL BE PROVIDED.**

Immediately examine your inventory and quarantine product subject to recall. In addition, if you may have further distributed these products, please identify your distributor and notify. Your notification to your distributor may be enhanced by including a copy of this recall notification letter.

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

Please complete and return the enclosed response form as soon as possible and return via one of the following methods.

Fax: 817-868-5362

Email: [rxrecalls@inmar.com](mailto:rxrecalls@inmar.com)

Address: Inmar Rx Solutions, Inc., Attn: Recall Coordinator – One West Fourth Street, Suite 500  
Winston Salem, NC 27101.

**If you have any questions about this event or the product return process, please contact Inmar Rx Solutions, Inc at 877-213-3809 between office hours of 9 AM to 5 PM EST, Monday through Friday.**

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

1. **Distributors/Pharmacies** – Immediately examine your inventory, quarantine and discontinue distribution of these lots.
2. **Distributors** – Complete the enclosed Business Response Form even if you do not have any product on hand.

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3. **Distributors** – Please pass this Recall Notice on **ONLY** to pharmacies that received these product lots.
4. **Pharmacies** – If you have units of the affected lots in inventory, please contact Inmar Rx Solutions, Inc. at 877-213-3809 to receive a Business Recall Response form or acquire it from <https://clsnetlink.com>.
5. Business Recall Response Form can be submitted by any of these methods.  
Fax: 817-868-5362  
Email: [rxrecalls@inmar.com](mailto:rxrecalls@inmar.com)  
Address: Inmar Rx Solutions, Inc., Attn: Recall Coordinator – One west fourth Street, Suite 500  
Winston Salem, NC 27101
6. **Distributors/Pharmacies** – Return the recalled product lots to Inmar Rx Solutions, Inc. as instructed in recall/return packet.
7. **Pharmacies** – You do not need to contact any patients.

**Upon receipt of the complete BRRF, a return kit will be sent including an RA form and necessary box labels.**

This recall should be carried out to the **retail** level.

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,

**Jigar Thakkar**  
Digitally signed by Jigar Thakkar  
Date: 2026.04.24 17:01:34 -04'00'

Jigar Thakkar  
Manager, Quality Assurance



# Lupin Pharmaceuticals, Inc.

## Label (s):

### a) Liraglutide Injection 6 mg/mL (2s Pack) carton label:

**Liraglutide Injection**  
18 mg/3 mL (6 mg/mL)

Use as directed by your healthcare professional.  
Subcutaneous use only  
Rx only

**Ingredients:** Each mL contains liraglutide 6 mg and the following inactive ingredients: disodium hydrogen phosphate dihydrate, 1.42 mg; phenol, 5.5 mg; propylene glycol, 14 mg; and water for injection.

**Sterile**  
**DO NOT TRANSFER THIS MEDICATION TO A SYRINGE**  
Each prefilled pen contains 18 mg/3 mL (6 mg/mL) and will deliver 36 doses of 0.6 mg or 15 doses of 1.2 mg or 10 doses of 1.8 mg.  
**REFRIGERATE – DO NOT FREEZE**  
After first use, store at a temperature not to exceed 86°F (30°C).  
Not a child-resistant container  
If the seal is broken before first use, contact your pharmacist.

M. L. No. MH/103164

Manufactured for:  
Lupin Pharmaceuticals, Inc.  
Naples, FL 34108  
United States  
Product of China

Manufactured by:  
Lupin Limited  
Nagpur 441106, INDIA.

**Liraglutide Injection**  
18 mg/3 mL (6 mg/mL)

2 Pens

NDC 70748-346-02

**Liraglutide Injection**  
18 mg/3 mL (6 mg/mL)

**For Single Patient Use Only**  
Each pen delivers doses of 0.6 mg, 1.2 mg or 1.8 mg.  
For subcutaneous use only  
Discard pen 30 days after first use  
**REFRIGERATE – DO NOT FREEZE**

Rx only  
LUPIN®

Contains: 2 Liraglutide Injection Pens, Product Literature  
**PHARMACIST:** Dispense the accompanying Medication Guide to each patient  
Pen needles not included

2 Pens

**Liraglutide Injection**  
18 mg/3 mL (6 mg/mL)

2 Pens

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# Lupin Pharmaceuticals, Inc.

b) Liraglutide Injection 6 mg/mL (3s Pack) carton label:

**Liraglutide Injection**  
**18 mg/3 mL (6 mg/mL)**  
 Use as directed by your healthcare professional.  
 See enclosed product literature.  
 Subcutaneous use only  
 Rx only

**Ingredients:** Each mL contains liraglutide 6 mg and the following inactive ingredients: disodium hydrogen phosphate dodecahydrate, 1.42 mg; phenol, 5.5 mg; propylene glycol, 14 mg; and water for injection.

**Sterile**  
**DO NOT TRANSFER THIS MEDICATION TO A SYRINGE**  
 Each prefilled pen contains 18 mg/3 mL (6 mg/mL) and will deliver 30 doses of 0.6 mg or 15 doses of 1.2 mg or 10 doses of 1.8 mg.

**REFRIGERATE - DO NOT FREEZE**  
 After first use, store at a temperature not to exceed 86°F (30°C).  
 Not a child-resistant container.  
 If the seal is broken before first use, contact your pharmacist.

M. L. No. MH103164

Manufactured by:  
 Lupin Pharmaceuticals, Inc.  
 Naples, FL 34108  
 United States  
 Product of China

Manufactured by:  
 Lupin Limited  
 Nagpur 441108, INDIA

**Liraglutide Injection**  
**18 mg/3 mL (6 mg/mL)**  
 3 Pens

**Liraglutide Injection**  
**18 mg/3 mL (6 mg/mL)**  
 NDC 70748-346-03  
**For Single Patient Use Only**  
 Each pen delivers doses of 0.6 mg, 1.2 mg or 1.8 mg.  
 For subcutaneous use only.  
 Discard pen 30 days after first use.  
**REFRIGERATE - DO NOT FREEZE**  
 Rx only  
**LUPIN®**

Contains: 3 Liraglutide Injection Pens, Product Literature  
 PHARMACIST: Dispense the accompanying  
 Medication Guide to each patient.  
 Pen needles not included.

**Liraglutide Injection**  
**18 mg/3 mL (6 mg/mL)**  
 3 Pens

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# Lupin Pharmaceuticals, Inc.

c) Liraglutide Injection 6 mg/mL prefilled pen label:

**NDC 70748-346-01**

**Liraglutide Injection**

**18 mg/3 mL (6 mg/mL)**

**For Single Patient Use Only**

Each prefilled pen contains 18 mg/3 mL (6 mg/mL) and will deliver 30 doses of 0.6 mg or 15 doses of 1.2 mg or 10 doses of 1.8 mg.

REFRIGERATE BEFORE FIRST USE – DO NOT FREEZE  
Discard pen 30 days after first use.

**Rx only**  
**LUPIN®**

Manufactured by:  
**Lupin Limited**  
Nagpur 441108, INDIA.  
M.L. No. MH/103164

276864

3 70748 34601 3

Lot .:

Exp.:

**Unvarnish Area**  
**37 x 15 mm**

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N131466 RCL096-26

### Lupin Pharmaceuticals, Inc.

#### RECALL

LIRAGLUTIDE Injection, 6MG/ML 3's

LIRAGLUTIDE Injection, 6MG/ML 2's

Retail Level

4/24/2026

**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 it is not provided, the processing of your form will be delayed.**

Address:

City:

State:

Zip:

Contact Name (Please Print):

Telephone#:

Email:

Contact Signature:

Date:

**DEBIT MEMO# (If unsure, leave blank):** \_\_\_\_\_

**Wholesaler Information if not directly purchased from Lupin:**

Wholesaler Name:

DEA#:

City:

State:

Zip:

**I have checked my stock and communicated to my customers at the appropriate level:**

I *confirm* that all locations that received the impacted products have been notified to the retail level. (Circle One) **YES** **YES-Corporate Notified** **NO (Why?)** \_\_\_\_\_

I do not have any stock of the recalled items. **OR**

I have quarantined and listed in the box below the quantity of recalled units and I will be returning to Inmar, as soon as possible. Upon receipt of this Response Form, Inmar, will issue return authorization label(s). Please indicate the # of needed box labels \_\_\_\_\_.

Product Name	NDC#	Lot#	Expiration Date	Total Full Syringes	Total Partial Syringes
LIRAGLUTIDE Injection, 6MG/ML 3's	70748-346-03	WB00098	7/31/2027		
LIRAGLUTIDE Injection, 6MG/ML 3's	70748-346-03	WB00092	7/31/2027		
LIRAGLUTIDE Injection, 6MG/ML 3's	70748-346-03	WB00093	7/31/2027		
LIRAGLUTIDE Injection, 6MG/ML 3's	70748-346-03	WB00104	10/31/2027		
LIRAGLUTIDE Injection, 6MG/ML 3's	70748-346-03	WB00106	10/31/2027		
LIRAGLUTIDE Injection, 6MG/ML 3's	70748-346-03	WB00105	10/31/2027		

See More NDC/Lots on Page 2

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Product Name	NDC#	Lot#	Expiration Date	Total Full Syringes	Total Partial Syringes
LIRAGLUTIDE Injection, 6MG/ML 2'S	70748-346-02	WB00097	7/31/2027		
LIRAGLUTIDE Injection, 6MG/ML 2'S	70748-346-02	WB00094	7/31/2027		
LIRAGLUTIDE Injection, 6MG/ML 2'S	70748-346-02	WB00088	7/31/2027		
LIRAGLUTIDE Injection, 6MG/ML 2'S	70748-346-02	WB00103	10/31/2027		
LIRAGLUTIDE Injection, 6MG/ML 2'S	70748-346-02	WC00016	12/31/2027		

If you have any questions regarding this form or product return please contact Inmar at 877-213-3809  
Office hours 9am to 5pm EST Mon thru Fri.

**Please fax this form to: 1-817-868-5362 or E-mail [rxrecalls@inmar.com](mailto:rxrecalls@inmar.com)**

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