# Gabapentin Capsules, USP, 100 mg Class II - Retail-Level Recall CUSTOMER CARE TALK TRACK

**Background Information:** NCQA made an update to their standard requesting communication be sent to members and prescribers affected by any recall classified as a Class II by the FDA except for wholesale level recalls. Communication is being sent to members and prescribers of NCQA delegated clients to inform them of the limited recall. **The distributor is not requiring any action on the part of consumers for these recalls**.

**This recall affects:**

|  |  |  |
| --- | --- | --- |
| **Product** | **NDC** | **Lot # / Exp. Date** |
| Gabapentin Capsules, USP, 100 mg, 100 capsules (10x10), blister pack cartons | 00904-6665-61 | M05205 exp. 10/2026 |
| Gabapentin Capsules, USP, 100 mg, 10 capsules (10x1) per bag | 55154-3363-00 | M05205A exp. 10/2026 |
| M05205B exp. 10/2026 |

# Customer Care Talk Track:

\*\* Please ensure to disposition all recall calls to code “**1116**” \*\*

|  |  |  |
| --- | --- | --- |
| **Q#** | **Question** | **Answer** |
| **1** | **Why did I receive a recall notice?** | Our records indicate that you may have recently received a prescription for a product affected by a limited recall from your retail pharmacy.  For more information, call the distributor listed on the letter or visit their website. You may also call the United States Food and Drug Administration (FDA) toll-free at 1‑888‑INFO-FDA (1‑888-463-6332) or visit [www.fda.gov](http://www.fda.gov). |
| **2** | **May I return the rest of the recalled product that I have?** | The distributor is not requiring any action on the part of consumers for these recalls. |
| **3** | **Should I stop using the recalled product?** | Please contact your prescriber with any questions or concerns about this recall or your use of the product. |

**Additional Information:**

**Event Details**

* **Event ID:** 97087
* **Voluntary / Mandated:** Voluntary: Firm initiated
* **Product Type:** Drugs
* **Initial Firm Notification of Consignee or Public:** Letter
* **Status:** Ongoing
* **Distribution Pattern:** Nationwide
* **Recalling Firm:**

The Harvard Drug Group LLC  
7000 Cardinal Pl  
Dublin, OH 43017-1091  
United States

* **Press Release URL(s):** Press Release Not Issued For This Recall
* **Recall Initiation Date:** 6/19/2025
* **Center Classification Date:** 7/3/2025
* **Date Terminated:** N/A
* **\*N/A -**Not Available

**Product Details**

| **#** | **Product Description** | **Recall Number** | **Classification** | **Code Information** | **Product Quantity** | **Reason for Recall** |
| --- | --- | --- | --- | --- | --- | --- |
| 1 | Gabapentin Capsules, USP, 100 mg, 100 capsules (10x10), blister pack cartons, Rx only, Packaged and Distributed by: MAJOR PHARMACEUTICALS, Indianapolis, IN, 46268 USA, NDC 0904-6665-61 | D-0507-2025 | Class II | Lot# M05205, Exp Date 10/2026 | 23,232 cartons | Defective container; blister packaging inadequately sealed. |
| 2 | Gabapentin Capsules, USP, 100 mg, 10 capsules (10x1) per bag, Rx only, Packaged and Distributed by: MAJOR PHARMACEUTICALS, Indianapolis, IN, 46268 USA, Distributed by Cardinal Health, Dublin, OH 43017, NDC 55154-3363-0 | D-0508-2025 | Class II | Lot# M05205A and M05205B, Exp Date 10/2026. | 3,527 bags | Defective container; blister packaging inadequately sealed. |

**Update History:** There is no history available for products in this event

[Top of the Document](#_top)

Not to Be Reproduced or Disclosed to Others without Prior Written Approval

**ELECTRONIC DATA = OFFICIAL VERSION / PAPER COPY = INFORMATIONAL ONLY**