

**RECALL RETURN RESPONSE FORM**  
**DILTIAZEM HYDROCHLORIDE EXTENDED RELEASE CAPSULES 120 MG (100's PACK**  
**CONTAINER)**  
**(NDC 68462-562-01)**  
**Retail Level**  
**04/16/2024**

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

Customer Name:		DEA#:
<i>DEA # is required, if it is not provided, the processing of your form will be delayed.</i>		
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 Glenmark Pharmaceuticals Inc.:**

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 \_\_\_\_\_ (Initial and date)

☐ 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\_\_\_\_\_.

Item Description	NDC#	Lot#/ Pack Size	Exp. Date	Total Full/Sealed and Partial/Open Bottle Count
DILTIAZEM HYDROCHLORIDE EXTENDED-RELEASE CAPSULES 120 MG	68462-562- 01	17221312/100's Pack Container	MAY- 2024	

If you have any questions regarding this form or product return please contact Inmar at 877-861-4963  
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)  
Recall Event ID N131161 / RCL110-24**