

RECALL RETURN RESPONSE FORM

Product Name : Hydrocodone Bitartrate and Acetaminophen Tablets, USP
Package Size : Bottle of 100's count
NDC Number : 31722-943-01
Lot # & Exp Date : 20070518 & June 2022

Please check all Appropriate Boxes

- ☐ I have read and understand the recall instructions provided in the notification dated 10/26/2020.
- ☐ I have checked my stock and have quarantined inventory consisting of _____ bottles/tablets.
- ☐ Indicate disposition of recalled product:
- ☐ Returned (**specify quantity, date and method** -----)/held for return-----.
- ☐ Destroyed (**specify quantity, date and method**) -----
- ☐ I have identified and notified my customers that were shipped or may have been shipped this product by (**specify date and method of notification**)

<or>

- ☐ Attached is a list of customers who received/ may have received this product. Please notify my customers.

Any adverse events associated with recalled product? ☐ Yes ☐ NO.

If yes, please explain: _____

Please check the appropriate box (es) to describe your business

☐ wholesaler/ distributor

☐ retailer

☐ hospital pharmacies

☐ hospital/medical facility

☐ pharmacy

☐ hospital pharmacies

☐ Others : _____

Name: _____

Title: _____

Phone: _____

Business Name: _____

Address: _____

DEA Number: _____

If Applicable:

Wholesaler Name: _____

Wholesaler DEA number: _____

If you have any questions regarding this form or product return, please contact Inmar at 855-607-9076.

Office hours 9am to 5pm EST Mon thru Fri.

Please fax this form to: 1-817-868-5362 or E-mail rxrecalls@inmar.com