RECALL STOCK RESPONSE FORM

Retail level Recall (ELIGARD® 7.5mg (leuprolide acetate) for injectable suspension)

VOLUNTARY RECALL – 12Sep2023

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.

Company 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_____

I have checked my stock and:

_____Do not have any stock of the recalled **items**.

OR

Have quarantined and listed in the box below the qty of recalled units 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 _____.

If Returning Pallets please indicate the number of pallets and the weight of each. _____pallet(s)_____weight

Email address for freight contact person _____

Drug	NDC	Lot	Exp.	Qty of product returning
ELIGARD® 7.5mg (leuprolide acetate) for injectable suspension, for subcutaneous use	**62935-753-75**	**13635A1**	**07 / 2024**	

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

Purchased From: Wholesaler Name _____DEA #_____

City_____ State_____

If you have any questions regarding this form or product return please contact Inmar at **1-855-296-8634** Office hours 9am to 5pm EST, Mon thru Fri.

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