



## **URGENT: DRUG RECALL – RESPONSE FORM** **(REVISED TO RETAIL LEVEL)**

**Please Complete This Form and Fax to: 817-868-5362**

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

Product Name	Package Description	Lot Number	NDC Number	Expiration Date
Medroxyprogesterone Acetate Injectable Suspension USP, 150 mg/ml	1 ml Pre-Filled Syringe	JKX4312A	50102-591-40	09/2022
		JKX4313A	50102-591-40	09/2022
		JKX4827A	50102-591-40	09/2023
		HAC1290A	50102-591-40	06/2023
		HAC2082B	50102-591-40	06/2023
		HAC1289A	16714-999-01	06/2023
		JKX2679A	16714-999-01	06/2022
		JKX3762A	16714-999-01	08/2022
		HAC0164A	16714-999-01	06/2023
		HAC1951A	62756-091-40	06/2023
	1 ml Vial	HAC2075A	16714-981-01	06/2023
		HAC2076A	16714-981-01	07/2023
		HAC2077A	16714-981-01	08/2023
		HAC2078A	16714-981-01	08/2023
		HAC3803A	16714-981-01	09/2023
		HAC0551A	16714-981-01	02/2023
		HAC0562A	16714-981-01	03/2023
		HAC1183A	16714-981-01	03/2023
		HAC1807A	16714-981-01	06/2023
		JKX6017A	16714-981-01	12/2022
		JKX6018A	16714-981-01	12/2022
		HAC0163A	16714-981-01	01/2023
		HAC1184A	16714-981-01	04/2023
		HAC0162A	16714-981-01	12/2022
		HAC2074A	62756-090-40	06/2023
		HAC0163B	62756-090-40	01/2023
		HAC1741A	62756-090-40	04/2023

For return of affected product, please email [rxrecalls@inmar.com](mailto:rxrecalls@inmar.com) or call 1- 855-884-7515.



## **URGENT: DRUG RECALL – RESPONSE FORM (REVISED)**

**Please Complete This Form and Fax to: 817-868-5362**  
**or Email to: [rxrecalls@inmar.com](mailto:rxrecalls@inmar.com)**

**Please check ALL appropriate boxes.**

☐ I have read and understand the recall instructions provided in the May 17, 2022 letter.

☐ I have checked our stock and have quarantined inventory consisting of \_\_\_\_\_ units.

☐ Indicate disposition of recalled product:

☐ returned (**specify quantity, date and method**)/held for return;

Number of Labels Required for Return to Inmar: \_\_\_\_\_

☐ previously destroyed (**specify quantity, date and method**);

☐ I have identified and notified my retail customers that were shipped or may have been shipped this product by (**specify date and method of notification**); or

☐ Attached is a list of retail 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: \_\_\_\_\_

For return of affected product, please email [rxrecalls@inmar.com](mailto:rxrecalls@inmar.com) or call 1-855-884-7515.



## **URGENT: DRUG RECALL – RESPONSE FORM (REVISED)**

**Please Complete This Form and Fax to: 817-868-5362**  
**or Email to: [rxrecalls@inmar.com](mailto:rxrecalls@inmar.com)**

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

- |                                                         |                                                    |
|---------------------------------------------------------|----------------------------------------------------|
| <input type="checkbox"/> wholesaler/distributor         | <input type="checkbox"/> retailer                  |
| <input type="checkbox"/> grocery corporate headquarters | <input type="checkbox"/> hospital pharmacies       |
| <input type="checkbox"/> repacker                       | <input type="checkbox"/> hospital/medical facility |
| <input type="checkbox"/> pharmacy                       | <input type="checkbox"/> Other:                    |

Customer Name: \_\_\_\_\_ Title: \_\_\_\_\_

Company: \_\_\_\_\_ DEA Number: \_\_\_\_\_

Address: \_\_\_\_\_

City: \_\_\_\_\_ State: \_\_\_\_\_ Zip Code: \_\_\_\_\_

Phone Number: \_\_\_\_\_

Customer Debit Memo Number: \_\_\_\_\_

Wholesaler: \_\_\_\_\_ City\State: \_\_\_\_\_

Wholesaler DEA Number: \_\_\_\_\_