

**RECALL STOCK RESPONSE FORM**

**RECALL of (AHP Pramipexole Dihydrochloride Tablets)  
(Retail Level)  
(05/13/2019)**

**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.

Customer Name \_\_\_\_\_ DEA # \_\_\_\_\_  
*\*DEA # is required, if it is 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**

I have quarantined and listed in the box below the quantity of recall 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 \_\_\_\_\_.

Product Description	AHP Lot No.	Expiration Date	Qty Returning
Pramipexole Dihydrochloride Tablets, 0.125 mg, 30 UD Carton NDC#: 68084-793-25 (Individual Dose NDC: 68084-793-95)	179049	12/31/2019	
	182571	8/31/2020	
Pramipexole Dihydrochloride Tablets, 0.25 mg, 100 UD Carton NDC#: 68084-440-01 (Individual Dose NDC: 68084-440-11)	172669	05/31/2019	
	175872	09/30/2019	
	177086	09/30/2019	
	179047	12/31/2019	
	182584	07/31/2020	
Pramipexole Dihydrochloride Tablets, 0.5 mg, 30 UD Carton NDC# :68084-974-25 (Individual Dose NDC: 68084-974-95)	175820	09/30/2019	
	176569	09/30/2019	
	177866	12/31/2019	
	179627A	12/31/2019	
	179627B	12/31/2019	
	181627	06/30/2020	
Pramipexole Dihydrochloride Tablets, 1 mg, 30 UD Carton NDC#: 68084-982-25 (Individual Dose NDC: 68084-982-95)	176179	05/31/2019	
	176616	07/31/2019	
	178562	09/30/2019	
	179947	12/31/2019	
	182048	04/30/2020	
	183136	06/30/2020	
	184217	08/31/2020	

**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 1-800-967-5952. 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)**