

URGENT DRUG RECALL- Retail Level BUSINESS RESPONSE FORM

12/23/2020

VUMERITY® 231 mg bottles

NDC#	LOT#	EXP DATE
101801 (106 ct)	64406-020-01	31-May-21
101799 (120 ct)	64406-020-03	31-May-21
102826 (106 ct)	64406-020-01	30-Jun-21
102362 (120 ct)	64406-020-03	30-Jun-21

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

Customer Name	DEA #_		
Address			
City	State	Zip	
Contact Name (please print)	Telepho	one #	
Fax #			
Contact Email			
Contact Signature			Date
I have read and understand the recall instruction I have identified and notified my customers.	·		
I have checked my stock and:			
[] Do not have any stock of the recalled items.			
OR			
[] I have quarantined and listed in the table be as possible. Upon receipt of this Response Form, product.	· · ·		-

Product Description	Lot Numbers	NDC	Sealed bottle quantity to be returned	Open bottle quantity to be returned
VUMERITY® 231 mg bottles	101801 (106 ct)	64406-020-01		
VUMERITY® 231 mg bottles	101799 (120 ct)	64406-020-03		
VUMERITY® 231 mg bottles	102826 (106 ct)	64406-020-01		
VUMERITY® 231 mg bottles	102362 (120 ct)	64406-020-03		

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

Purchased From: Wholesaler Name		Wholesaler DEA#		
Any adverse events associated with recalled/failed product?	No []	Yes [] If yes, please explain:		

Please notify your direct consignees to the **RETAIL** level

If you have any questions regarding this form or product return please contact Inmar Customner Service (1-800-967-5952 during the hours of 9am to 5pm EST, Monday through Friday.

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