

## URGENT DRUG RECALL BUSINESS RESPONSE FORM

## 12/10/2020

PRODUCT DESCRIPTION	NDC#	LOT#	EXP DATE
Azacitidinefor Injection 100mg/vial	69097-805-40	7S10182A	Sep 2021

<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				
Contact Name (please print)		Telephone #		
Fax #				
Contact Email				
Contact Signature			Date	
[ ] I have read and understand the r	ecall instructions provid	ed in the letter.		
[ ] I have identified and notified my	customers that were sh	ipped this product.		
I have checked my stock and:				
[ ] Do not have any stock of the rec	alled items.			
OR				
[ ] I have quarantined and listed in a soon as possible. Upon receipt of this with the product.		•	_	

Product Description	NDC	Lot Numbers	Sealed bottle quantity to be returned	Open bottle quantity to be returned
Azacitidine for Injection 100 mg/vial	69097-805-40	7S10182A		

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:		
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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: <a href="mailto:rxrecalls@inmar.com">rxrecalls@inmar.com</a>