




SAGENT®

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CUSTOMER RECALL RETURN RESPONSE FORM

PLEASE FAX COMPLETED RESPONSE FORM TO 1-901-368-6903 or EMAIL TO ddnregulatory@eversana.com

Product	Lot Number	Expiration Date	NDC Number	Distribution Dates
Ketorolac Tromethamine Injection, USP 	M813513	Feb-2020	25021-701-02	January - March 2019

Please check ALL appropriate boxes.

- I have read and understand the recall instructions provided in the Customer Notification/Recall Communication letter dated April 30, 2019.
- I have checked my stock and have quarantined inventory consisting of _____ units.

Indicate disposition of recalled product:

Disposition	Lot	Quantity	Date	Method
<input type="checkbox"/> returned				
<input type="checkbox"/> destroyed				
<input type="checkbox"/> relabeled				
<input type="checkbox"/> quarantined				

I have identified and notified my customers that were shipped or may have been shipped this product and have communicated that we are conducting a sub-recall to our direct account customers. This recall should be carried out to the user level.

Have there been any Adverse Events associated with recalled product? Yes NO

If yes, please explain: _____

Please check the appropriate box(es) to describe your business			
<input type="checkbox"/> wholesaler/distributor	<input type="checkbox"/> retailer	<input type="checkbox"/> pharmacy – retail	<input type="checkbox"/> hospital pharmacies
<input type="checkbox"/> hospital/medical facility	<input type="checkbox"/> medical laboratory	<input type="checkbox"/> Other:	

Please Complete Contact Information for Person Completing Response:	
Name:	
Title:	
Tel Number:	
Facility:	
Address:	
City, State, Zip:	