

FDA
DEPARTMENT OF HEALTH AND HUMAN SERVICES
 Food and Drug Administration

MEDWATCH
FORM 3500A

 For use by user-facilities, importers, distributors
 and manufacturers for MANDATORY reporting

Form Approved: OMB No. 0910-0291

Expires: 6-30-2025

See PRA statement on page 6.

FDA USE ONLY

Mfr report #

UF/Importer Report #

Exemption/Variance #

Note: For date prompts of "dd-mmm-yyyy" please use 2-digit day, 3-letter month abbreviation, and 4-digit year; for example, 01-JAN-1900.
A. PATIENT INFORMATION

1. Patient Identifier (<i>In confidence</i>)		2. Age		or Date of Birth (<i>e.g., 01-Jan-1900</i>)	
		Year(s)	Week(s)		
		Month(s)	Day(s)		
3. Sex: Enter the patient's sex at birth (<i>the sex that a person has or was assigned to at birth</i>). Male Female		SECTION REMOVED			
4. Weight lb kg	5. Ethnicity (<i>Check one</i>) Hispanic/Latino Not Hispanic/Latino	6. Race (<i>check all that apply</i>) American Indian/Alaska Native Native Hawaiian/ Asian Other Pacific Islander Black or African American White			

B. ADVERSE EVENT OR PRODUCT PROBLEM

1. Type of Report (<i>check all that apply</i>) Adverse Event Product Problem (<i>e.g., defects/malfunctions</i>)		2. Outcome Attributed to Adverse Event (<i>check all that apply</i>) Death – Date of death (<i>01-JAN-1900</i>): Life-threatening Hospitalization (initial or prolonged) Other Serious or Important Medical Events Required Intervention to Prevent Permanent Impairment/Damage Disability or Permanent Damage Congenital Anomaly/Birth Defects	
3. Date of Event (<i>01-JAN-1900</i>)	4. Date of this Report (<i>01-JAN-1900</i>)		

Submission of a report does not constitute an admission that medical personnel or the product caused or contributed to the event.

* Please see instructions

5. Describe Event or Problem

6. Relevant Test/Laboratory Data

Date (01-JAN-1900)

Relevant Test/Laboratory Data

Date (01-JAN-1900)

Additional comments

7. **Other Relevant History, Including Preexisting Medical Conditions** (e.g., allergies, pregnancy, tobacco product use, alcohol use, and liver/kidney problems, etc.)

C. SUSPECT PRODUCTS

SUSPECT PRODUCT #1

1. Name, Strength, Manufacturer/Compounder

Product Name		Strength	Unit
NDC # or Unique ID	Manufacturer/Compounder Name		Lot #

2. List Medical Product and Treatment Given at the Same Time of the Event and Date (Do not include treatment for initial event)

3. Dose or Amount	Frequency	Route
Unit	Other Frequency	Other Route

4. Treatment Dates/Therapy Dates (give best estimate of length of treatment (start/stop) or date of dose reduction.)

Therapy started on (e.g., 01-Jan-1900)	Therapy stopped on (e.g., 01-Jan-1900)	Dose Reduced (e.g., 01-Jan-1900)	OR	Duration	Unit
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5. Diagnosis for use (indication)

6. Product Type (check all that apply)

7. Expiration Date (e.g., 01-Jan-1900)

OTC		Generic	
Compounded		Biosimilar	

8. Event Abated after use Stopped or Dose Reduced?

9. Event Reappeared after Reintroduction?

Yes No Doesn't apply

Yes No Doesn't apply

SUSPECT PRODUCT #2**1. Name, Strength, Manufacturer/Compounder**

Product Name	Strength	Unit
NDC # or Unique ID	Manufacturer/Compounder Name	Lot #

2. List Medical Product and Treatment Given at the Same Time of the Event and Date *(Do not include treatment for initial event)***3. Dose or Amount****Frequency****Route**

Unit

Other Frequency

Other Route

4. Treatment Dates/Therapy Dates *(give best estimate of length of treatment (start/stop) or date of dose reduction.)*

Therapy started on <i>(e.g., 01-Jan-1900)</i>	Therapy stopped on <i>(e.g., 01-Jan-1900)</i>	Dose Reduced <i>(e.g., 01-Jan-1900)</i>	OR	Duration	Unit
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5. Diagnosis for use *(indication)***6. Product Type** *(check all that apply)***7. Expiration Date** *(e.g., 01-Jan-1900)*

OTC

Generic

Compounded

Biosimilar

8. Event Abated after use Stopped or Dose Reduced?

Yes

No

Doesn't apply

9. Event Reappeared after Reintroduction?

Yes

No

Doesn't apply

D. SUSPECT MEDICAL DEVICE**1. Brand Name****2a. Common Device Name****2b. Procode****3. Manufacturer Name, City and State****4. Model #****Lot #****Catalog #****Expiration Date** *(01-JAN-1900)***Serial #**

Unique Device Identifier (UDI) #																																				
5. Operator of Device Health Professional Patient/Consumer Other		6a. If Implanted, Give Date (01-JAN-1900)	6b. If Explanted, Give Date (01-JAN-1900)																																	
7a. Is this a single-use device that was reprocessed and reused on a patient? Yes No		7b. If yes, enter the name and address of the reprocessor																																		
8. Was this device ever serviced by a third-party servicer? Yes No Unknown		9. Is this Device Available for Evaluation? (Do not send to FDA) Yes No Returned to manufacturer on (01-JAN-1900)																																		
10. Concomitant Medical Products and Therapy Dates (Exclude treatment of event) <table border="1" style="width: 100%; border-collapse: collapse; margin-top: 5px;"> <thead> <tr> <th style="width: 45%;">Product Name</th> <th style="width: 25%;">Therapy Start Date (01-JAN-1900)</th> <th style="width: 30%;">Therapy End Date (01-JAN-1900)</th> </tr> </thead> <tbody> <tr><td>1.</td><td></td><td></td></tr> <tr><td>2.</td><td></td><td></td></tr> <tr><td>3.</td><td></td><td></td></tr> <tr><td>4.</td><td></td><td></td></tr> <tr><td>5.</td><td></td><td></td></tr> <tr><td>6.</td><td></td><td></td></tr> <tr><td>7.</td><td></td><td></td></tr> <tr><td>8.</td><td></td><td></td></tr> <tr><td>9.</td><td></td><td></td></tr> <tr><td>10.</td><td></td><td></td></tr> </tbody> </table>				Product Name	Therapy Start Date (01-JAN-1900)	Therapy End Date (01-JAN-1900)	1.			2.			3.			4.			5.			6.			7.			8.			9.			10.		
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1. Name and Address <table border="1" style="width: 100%; border-collapse: collapse; margin-top: 5px;"> <tr> <td colspan="2" style="height: 30px; vertical-align: bottom;">Last Name</td> <td colspan="2" style="height: 30px; vertical-align: bottom;">First Name</td> </tr> <tr> <td colspan="4" style="height: 30px; vertical-align: bottom;">Address</td> </tr> <tr> <td style="width: 30%; height: 30px; vertical-align: bottom;">City</td> <td style="width: 20%; height: 30px; vertical-align: bottom;">State/Province/Region</td> <td style="width: 20%; height: 30px; vertical-align: bottom;">ZIP/Postal Code</td> <td style="width: 30%; height: 30px; vertical-align: bottom;">Country</td> </tr> <tr> <td colspan="2" style="height: 30px; vertical-align: bottom;">Phone #</td> <td colspan="2" style="height: 30px; vertical-align: bottom;">Email</td> </tr> </table>				Last Name		First Name		Address				City	State/Province/Region	ZIP/Postal Code	Country	Phone #		Email																		
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Phone #		Email																																		
2. Health Professional? Yes No		3. Occupation (Select from list)																																		
4. Initial reporter also sent report to FDA Yes No Unknown																																				

F. FOR USE BY USER FACILITY/IMPORTER (Devices Only)

1. Check One User Facility Importer		2. User Facility/Importer Report Number											
3. User Facility or Importer Name/Address		4. Contact Person		5. Phone Number									
		6. Date User Facility or Importer Became Aware of Event (01-JAN-1900)		7. Type of Report									
8. Date of This Report (01-JAN-1900)		9. Approximate Age of Device											
10. Adverse Event Problem (Refer to coding manual) <table style="width: 100%; border: none;"> <tr> <td style="width: 25%; border: none;">Health Effect – Clinical Code</td> <td style="width: 25%; border: none;">Health Effect – Impact Code</td> <td style="width: 25%; border: none;">Medical Device Problem Code</td> <td style="width: 25%; border: none;">Component Code</td> </tr> </table>					Health Effect – Clinical Code	Health Effect – Impact Code	Medical Device Problem Code	Component Code					
Health Effect – Clinical Code	Health Effect – Impact Code	Medical Device Problem Code	Component Code										
11. Report Sent to FDA? <i>(If Yes, enter date (01-JAN-1900))</i> Yes No		12. Location Where Event Occurred <table style="width: 100%; border: none;"> <tr> <td style="width: 25%; border: none;">Ambulatory Surgical Facility</td> <td style="width: 25%; border: none;">Outpatient Treatment Facility</td> <td style="width: 50%; border: none;">Other (Specify)</td> </tr> <tr> <td style="border: none;">Home</td> <td style="border: none;">Outpatient Diagnostic Facility</td> <td style="border: none;"></td> </tr> <tr> <td style="border: none;">Hospital</td> <td style="border: none;">Nursing Home</td> <td style="border: none;"></td> </tr> </table>			Ambulatory Surgical Facility	Outpatient Treatment Facility	Other (Specify)	Home	Outpatient Diagnostic Facility		Hospital	Nursing Home	
Ambulatory Surgical Facility	Outpatient Treatment Facility	Other (Specify)											
Home	Outpatient Diagnostic Facility												
Hospital	Nursing Home												
13. Report Sent to Manufacturer? <i>(If Yes, enter date (01-JAN-1900))</i> Yes No		14. Manufacturer Name/Address											

G. ALL MANUFACTURERS

1. Contact Office (and Manufacturing Site for Devices) or Compounding Outsourcing Facility <table style="width: 100%; border: none;"> <tr> <td style="width: 40%; border: none;">Name</td> <td style="width: 30%; border: none;">Email Address</td> <td style="width: 30%; border: none;">Phone Number</td> </tr> </table> Address					Name	Email Address	Phone Number						
Name	Email Address	Phone Number											
Compounding Outsourcing Facility 503B? Check box if applicable		Outsourcing Facility											
2. Report Source (check all that apply) <table style="width: 100%; border: none;"> <tr> <td style="width: 15%;">Foreign</td> <td style="width: 15%;">Literature</td> <td style="width: 15%;">Health Professional</td> <td style="width: 15%;">Company Representative</td> <td rowspan="2" style="width: 40%; vertical-align: bottom;">Other (Please list)</td> </tr> <tr> <td>Study</td> <td>Consumer</td> <td>Use Facility</td> <td>Distributor/Importer</td> </tr> </table>				Foreign	Literature	Health Professional	Company Representative	Other (Please list)	Study	Consumer	Use Facility	Distributor/Importer	3. Date Received by Manufacturer (01-JAN-1900)
Foreign	Literature	Health Professional	Company Representative	Other (Please list)									
Study	Consumer	Use Facility	Distributor/Importer										
4. NDA #	ANDA #	IND #	BLA #	PMA/510(k) #									
Check all that apply: Combination product Pre-ANDA Pre-1938 OTC Compounded Product													
5. If IND/Pre-ANDA, Give Protocol #		6. Type of Report (Check all that apply) <table style="width: 100%; border: none;"> <tr> <td style="width: 15%;">5-day</td> <td style="width: 15%;">15-day</td> <td style="width: 15%;">Periodic</td> <td style="width: 55%;">Follow-up #</td> </tr> <tr> <td>7-day</td> <td>30-day</td> <td>Initial</td> <td></td> </tr> </table>			5-day	15-day	Periodic	Follow-up #	7-day	30-day	Initial		
5-day	15-day	Periodic	Follow-up #										
7-day	30-day	Initial											
7. Adverse Event Term(s)				8. Manufacturer Report Number									

H. DEVICE MANUFACTURERS ONLY

1. Type of Reportable Event <i>(check all that apply.)</i> Death Malfunction Serious Injury Summary Report No. of events summarized		2. If Follow-up, What Type? Correction Additional Information Response to FDA Request Device Evaluation		3. Device Evaluated by Manufacturer? Yes No	
4. Device Manufacture Date (01-JAN-1900)			5. Labeled for Single Use? Yes No		
6. Adverse Event Problem <i>(Refer to coding manual)</i>					
Health Effect – Clinical Code		Health Effect – Impact Code		Medical Device Problem Code	
Type of Investigation		Investigation Findings		Investigation Conclusions	
7. If Remedial Action Initiated, Check Type Recall Relabeling Patient Monitoring Repair Notification Modification/Adjustment Replace Inspection Other:					8. Usage of Device Initial Use of Device Reuse Unknown
9. If action reported to FDA under 21 USC 360i(g), list correction/ removal reporting number:			10. Related Report Number		
11. Additional Manufacturer Narrative					

This section applies only to requirements of the Paperwork Reduction Act of 1995. This section applies only to requirements of the Paperwork Reduction Act of 1995. The public reporting burden for this collection of information has been estimated to average 73 minutes per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to:

Department of Health and Human Services
 Food and Drug Administration
 Office of Chief Information Officer
 Paperwork Reduction Act (PRA) Staff
 PRAStaff@fda.hhs.gov

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