

# SUSPECT ADVERSE REACTION REPORT

## I. REACTION INFORMATION

1. PATIENT INITIALS (first, last) <b>PRIVACY</b>	1a. COUNTRY <b>DOMINICAN REPUBLIC</b>	2. DATE OF BIRTH			2a. AGE <b>11 Years</b>	3. SEX <b>Female</b>	3a. WEIGHT <b>Unk</b>	4-6 REACTION ONSET			8-12 CHECK ALL APPROPRIATE TO ADVERSE REACTION  <input type="checkbox"/> PATIENT DIED  <input type="checkbox"/> INVOLVED OR PROLONGED INPATIENT HOSPITALISATION  <input type="checkbox"/> INVOLVED PERSISTENT OR SIGNIFICANT DISABILITY OR INCAPACITY  <input type="checkbox"/> LIFE THREATENING
		Day	Month	Year				Day	Month	Year	
		<b>PRIVACY</b>						<b>Unk</b>			

7 + 13 DESCRIBE REACTION(S) (including relevant tests/lab data)  
Event Verbatim [LOWER LEVEL TERM] (Related symptoms if any separated by commas)  
he device leaked medication when first inserted and the dosage was incorrect [Device leakage]  
the patient was supposed to get a dose of 10, but when she was turning the vial of the device, she realized that it only showed up to 3. [Inaccurate delivery by device]  
she had a bad practice in administering the medication [Wrong technique in device usage process]  
when first inserted and the dosage was incorrect [Incorrect dose administered by device]

(Continued on Additional Information Page)

## II. SUSPECT DRUG(S) INFORMATION

14. SUSPECT DRUG(S) (include generic name) #1 ) Genotropin Pen (SOMATROPIN) Solution for injection #2 ) Genotropin Pen (SOMATROPIN (DEVICE CONSTITUENT)) Solution for injection		20. DID REACTION ABATE AFTER STOPPING DRUG?  <input type="checkbox"/> YES <input type="checkbox"/> NO <input checked="" type="checkbox"/> NA
15. DAILY DOSE(S) #1 ) 1 mg, daily(at night) #2 )	16. ROUTE(S) OF ADMINISTRATION #1 ) Unknown #2 ) Unknown	
17. INDICATION(S) FOR USE #1 ) Unknown #2 ) Unknown		21. DID REACTION REAPPEAR AFTER REINTRODUCTION?  <input type="checkbox"/> YES <input type="checkbox"/> NO <input checked="" type="checkbox"/> NA
18. THERAPY DATES(from/to) #1 ) Unknown #2 ) Unknown	19. THERAPY DURATION #1 ) Unknown #2 ) Unknown	

## III. CONCOMITANT DRUG(S) AND HISTORY

22. CONCOMITANT DRUG(S) AND DATES OF ADMINISTRATION (exclude those used to treat reaction)		
23. OTHER RELEVANT HISTORY. (e.g. diagnostics, allergies, pregnancy with last month of period, etc.) From/To Dates                      Type of History / Notes                      Description Unknown		

## IV. MANUFACTURER INFORMATION

24a. NAME AND ADDRESS OF MANUFACTURER Pfizer S.A. Laura Arce Mora Avenida Escazú, Torre Lexus, piso 7. Escazú San Jose, COSTA RICA		26. REMARKS
	24b. MFR CONTROL NO. <b>PV202500086110</b>	
24c. DATE RECEIVED BY MANUFACTURER <b>27-AUG-2025</b>	24d. REPORT SOURCE <input type="checkbox"/> STUDY <input type="checkbox"/> LITERATURE <input type="checkbox"/> HEALTH PROFESSIONAL <input checked="" type="checkbox"/> OTHER: Spontaneous	
DATE OF THIS REPORT <b>27-AUG-2025</b>	25a. REPORT TYPE <input type="checkbox"/> INITIAL <input checked="" type="checkbox"/> FOLLOWUP: 3	
		25b. NAME AND ADDRESS OF REPORTER NAME AND ADDRESS WITHHELD.  NAME AND ADDRESS WITHHELD.  NAME AND ADDRESS WITHHELD.

27-Aug-2025 04:35

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**ADDITIONAL INFORMATION****7+13. DESCRIBE REACTION(S) continued**

Case Description: This is a spontaneous report and received from Consumer or other non HCPs from product quality group, Program ID: 164974.

An 11-year-old female patient received somatropin (GENOTROPIN PEN), (Batch/Lot number: unknown) at 1 mg daily (1 mg, daily(at night)). The patient's relevant medical history and concomitant medications were not reported.

The following information was reported: DEVICE LEAKAGE (non-serious), described as "he device leaked medication when first inserted and the dosage was incorrect"; DEVICE DELIVERY SYSTEM ISSUE (non-serious), described as "the patient was supposed to get a dose of 10, but when she was turning the vial of the device, she realized that it only showed up to 3."; WRONG TECHNIQUE IN DEVICE USAGE PROCESS (non-serious), described as "she had a bad practice in administering the medication"; INCORRECT DOSE ADMINISTERED BY DEVICE (non-serious), described as "when first inserted and the dosage was incorrect". The action taken for somatropin was unknown.

Product Quality Group provided investigational summary and conclusion on 29Jul2025 for somatropin (device constituent): Site Investigation (PUURS): No further investigation was required as no valid lot number or returned sample was available. This complaint will continue to be trended. If additional information becomes available, this complaint will be reopened. Device Investigation: This investigation is based on the information captured in the Complaint Description and Argus Report. The Complaint Issue, Leaking During Prep/Use, was reported. The Risk Management File was reviewed to confirm that the Hazard(s) and Hazardous Situation(s) associated with the Complaint Issue are documented in the Hazard Analysis (INX#100281795, Version # (9.0)). All complaint investigations are trended. There is no current trend alert documented.

Additional information: The patient's mother indicated that on the first day of the injection the medication spilled a little. The main problem occurred on the fifth day, when she went to administer the dose, she explained that the patient was supposed to get a dose of 10, but when she was turning the vial of the device, she realized that it only showed up to 3. Her consultation was to know what he should do in that situation. Reporter indicated that: "She was asking a question to the previous lady, since she had a bad practice in administering the medication." Reporter indicated that the device leaked medication when first inserted and the dosage was incorrect.

Causality for "he device leaked medication when first inserted and the dosage was incorrect", "the patient was supposed to get a dose of 10, but when she was turning the vial of the device, she realized that it only showed up to 3.", "she had a bad practice in administering the medication" and "when first inserted and the dosage was incorrect" was determined associated to device constituent of somatropin (malfunction).

Follow-up(29Jul2025): This is a follow-up report from product quality group providing investigation results. Updated information: Investigation results updated.

Follow-up (07Aug2025): This is a follow-up report from product quality group providing investigation results. Updated information includes investigation codes updated.

Follow-up (27Aug2025): Follow-up attempts are completed. Batch/lot number is not provided, and it cannot be obtained.