

SUSPECT ADVERSE REACTION REPORT

I. REACTION INFORMATION

1. PATIENT INITIALS (first, last) PRIVACY	1a. COUNTRY COSTA RICA	2. DATE OF BIRTH			2a. AGE 46 Years	3. SEX Female	3a. WEIGHT 68.00 kg	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 <input type="checkbox"/> CONGENITAL ANOMALY <input checked="" type="checkbox"/> OTHER
		Day	Month	Year				Day	Month	Year	
7 + 13 DESCRIBE REACTION(S) (including relevant tests/lab data) Event Verbatim [PREFERRED TERM] (Related symptoms if any separated by commas) Other Serious Criteria: Medically Significant Erosive gastritis [Gastritis erosive] Gained weight [Weight increased] Case Description: Study ID: 828652-My Healthy Journey Study description: Trial title: This is a 40 weeks digital patient support program with focus on exercise, motivation, nutrition & maintaining strategies (only for patients under Liraglutide 3.0 mg). (Continued on Additional Information Page)											

II. SUSPECT DRUG(S) INFORMATION

14. SUSPECT DRUG(S) (include generic name) #1) Saxenda (liraglutide 6 mg/mL) Solution for injection, 6 mg/mL {Lot # PP5M440; Exp.Dt. AUG-2026}		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) 0.60 mg, qd	16. ROUTE(S) OF ADMINISTRATION #1) Subcutaneous	
17. INDICATION(S) FOR USE #1) Weight loss (Weight control)		21. DID REACTION REAPPEAR AFTER REINTRODUCTION? <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> NA
18. THERAPY DATES(from/to) #1) 18-MAY-2025 / 23-MAY-2025	19. THERAPY DURATION #1) 5 days	

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 Novo Nordisk A/S Lise Grimmeshave Vandtaarnsvej 114 Soeborg, DK-2860 DENMARK Phone: +45 44448888		26. REMARKS Medically Confirmed: No
	24b. MFR CONTROL NO. 1454139	25b. NAME AND ADDRESS OF REPORTER NAME AND ADDRESS WITHHELD.
24c. DATE RECEIVED BY MANUFACTURER 09-JUN-2025	24d. REPORT SOURCE <input checked="" type="checkbox"/> STUDY <input type="checkbox"/> LITERATURE <input type="checkbox"/> HEALTH PROFESSIONAL <input type="checkbox"/> OTHER:	
DATE OF THIS REPORT 16-JUN-2025	25a. REPORT TYPE <input checked="" type="checkbox"/> INITIAL <input type="checkbox"/> FOLLOWUP:	

16-Jun-2025 13:18

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

Patient's height: 163 cm.
Patient's weight: 68 kg.
Patient's BMI: 25.59373710.

This serious Solicited Report from COSTA RICA was reported by a Consumer as "Erosive gastritis(Gastritis erosive)" beginning on 21-MAY-2025 , "Gained weight(Weight gain)" beginning on 21-MAY-2025 and concerned a 46 Years old Female patient who was treated with Saxenda (liraglutide 6 mg/mL) from 18-MAY-2025 to 23-MAY-2025 for "Weight loss",

Dosage Regimens:
Saxenda: 18-MAY-2025 to 23-MAY-2025;

Medical history was not provided.

Treatment medications included - RILATEN(ROCIVERINE), NEXIUM ESOMEPRAZOLE SODIUM, NEWDISPET(LEVOSULPIRIDE), LISALGIL(METAMIZOLE MAGNESIUM), SERTAL COMPUESTO FORTE(CLONIXIN LYSINATE, PARGEVERINE HYDROCHLORIDE).

Patient had the first session with the nutritionist who told patient to start as she indicated, but patient felt fine on the first day, and by the third day body was different.
Patient used a syringe for 5 days and have no underlying conditions.

On 21-MAY-2025, Patient began to vomit day and night, diarrhea, constant burping; nothing would go away and was burping like rotting.

On 21-MAY-2025, Patient experienced erosive gastritis and patient felt like was going to die and stomach swelled so much, and patient also gained Weight (values and units not reported).

Patient went to observation as symptoms wouldn't go away.

Patient took Rilaten and Lisalgil intravenously along with Sertal compuesto for pain. Patient took Nexium for gastritis and Newdispet for burping.

Batch Numbers:
Saxenda: PP5M440;

Action taken to Saxenda was reported as Product discontinued due to AE.

The outcome for the event "Erosive gastritis(Gastritis erosive)" was Recovering/resolving.
The outcome for the event "Gained weight(Weight gain)" was Recovering/resolving.

Reporter's causality (Saxenda) -
Erosive gastritis(Gastritis erosive) : Possible
Gained weight(Weight gain) : Possible

Company's causality (Saxenda) -
Erosive gastritis(Gastritis erosive) : Possible
Gained weight(Weight gain) : Unlikely

13. Lab Data

#	Date	Test / Assessment / Notes	Results	Normal High / Low
1	21-MAY-2025	Weight		
		Gained(Values and units not reported)		

13. Relevant Tests

On an unknown date, Patient gained weight (Values and units not reported)