

SUSPECT ADVERSE REACTION REPORT

I. REACTION INFORMATION

1. PATIENT INITIALS (first, last) PRIVACY	1a. COUNTRY COSTA RICA	2. DATE OF BIRTH			2a. AGE 58 Years	3. SEX Female	3a. WEIGHT 108.00 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 type="checkbox"/> OTHER
		Day	Month	Year			Day	Month	Year		
			PRIVACY					MAR	2025		

7 + 13 DESCRIBE REACTION(S) (including relevant tests/lab data)
Event Verbatim [PREFERRED TERM] (Related symptoms if any separated by commas)
 thyroid was very out of balance (Hypothyroidism aggravated) [Hypothyroidism] ([Condition aggravated])
 Gastritis (pain in the upper part of the stomach), bad-tasting belching, feel unwell [Gastritis]
 Nausea [Nausea]
 Vomiting [Vomiting]
 A loss of appetite [Decreased appetite]
 disgusted by food [Food aversion]
 Ozempic 0.5 mg applied in 18 clicks [Wrong technique in product usage process]

(Continued on Additional Information Page)

II. SUSPECT DRUG(S) INFORMATION

14. SUSPECT DRUG(S) (include generic name) #1) Semaglutide B 1.34 mg/ml PDS290 1.0 mg (SEAGLUTIDE 1.34 mg/mL) Solution for injection, 1 mg {Lot # (Continued on Additional Information Page)		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) 18 clicks to get 0.5 mg	16. ROUTE(S) OF ADMINISTRATION #1) Subcutaneous	
17. INDICATION(S) FOR USE #1) type 2 diabetes mellitus (Type 2 diabetes mellitus) (Continued on Additional Information Page)		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) 26-MAR-2025 / Unknown	19. THERAPY DURATION #1) Unknown	

III. CONCOMITANT DRUG(S) AND HISTORY

22. CONCOMITANT DRUG(S) AND DATES OF ADMINISTRATION (exclude those used to treat reaction) #1) EUTIROX (LEVOTHYROXINE SODIUM) ; Ongoing #2) NPH INSULIN (NPH INSULIN) ; Ongoing #3) INSULIN NOS (INSULIN NOS) ; Ongoing #4) XIGDUO (DAPAGLIFLOZIN PROPANEDIOL MONOHYDRATE, METF #5) VENLAFAXINE (VENLAFAXINE) Tablet ; 2023 / Ongoing #6) Novolin R Penfill (INSULIN HUMAN 100 IU/mL) Solution for injecti (Continued on Additional Information Page)		
23. OTHER RELEVANT HISTORY. (e.g. diagnostics, allergies, pregnancy with last month of period, etc.)		
From/To Dates	Type of History / Notes	Description
Unknown to Ongoing	Current Condition	Obesity (Obesity)
Unknown to Ongoing	Current Condition	Type 2 diabetes mellitus (Type 2 diabetes mellitus)

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. 1400838	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 24-JUN-2025	25a. REPORT TYPE <input checked="" type="checkbox"/> INITIAL <input type="checkbox"/> FOLLOWUP:	

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

Case Description: ***This is an auto generated narrative***

Study ID: 199-NovoDia

Study description: Trial Title: Patient support programme to support physician and their daily work to maintain an optimal diabetic control of patients through added value services such as treatment starter kit, nutrition support through NovoDia call center, individual workshops, group workshops and free A1c test.

Patient's height: 159 cm.

Patient's weight: 108 kg.

Patient's BMI: 42.71982910.

This non-serious Solicited Report from COSTA RICA was reported by a Consumer as "thyroid was very out of balance (Hypothyroidism aggravated)(Hypothyroidism)" beginning on 09-MAY-2025 , "thyroid was very out of balance (Hypothyroidism aggravated)(Condition aggravated)" beginning on 09-MAY-2025 , "Gastritis (pain in the upper part of the stomach), bad-tasting belching,feel unwell(Gastritis)" beginning on MAR-2025 , "Nausea(Nausea)" beginning on 26-MAR-2025 , "Vomiting(Vomiting)" beginning on 26-MAR-2025 , "A loss of appetite(Appetite lost)" beginning on 26-MAR-2025 , "disgusted by food(Food disgust)" beginning on 26-MAR-2025 , "Ozempic 0.5 mg applied in 18 clicks(Wrong technique in product usage process)" beginning on 26-MAR-2025 and concerned a 58 Years old Female patient who was treated with Ozempic 1.0 mg (SEMAGLUTIDE 1.34 mg/mL) from 26-MAR-2025 and ongoing for "type 2 diabetes mellitus", "Obesity",

Dosage Regimens:

Ozempic 1.0 mg: 26-MAR-2025 to Not Reported, ??-JUN-2025 to Not Reported (Dosage Regimen Ongoing);

Current Condition: Obesity, Type 2 diabetes mellitus, Hypothyroidism, Depression, Anxiety, High blood pressure.

Concomitant medications included - EUTIROX(LEVOTHYROXINE SODIUM), NPH INSULIN, INSULIN NOS, XIGDUO(DAPAGLIFLOZIN PROPANEDIOL MONOHYDRATE, METFORMIN HYDROCHLORIDE), VENLAFAXINE, Novolin R Penfill(INSULIN HUMAN 100 IU/mL), Novolin N(INSULIN ISOPHANE), BIOSARTAN(LOSARTAN POTASSIUM).

Batch Numbers:

Ozempic 1.0 mg: PP5L760, PP5L760;

Action taken to Ozempic 1.0 mg was reported as No Change.

The outcome for the event "thyroid was very out of balance (Hypothyroidism aggravated)(Hypothyroidism)" was Not recovered.
The outcome for the event "thyroid was very out of balance (Hypothyroidism aggravated)(Condition aggravated)" was Not recovered.

The outcome for the event "Gastritis (pain in the upper part of the stomach), bad-tasting belching,feel unwell(Gastritis)" was Recovering/resolving.

The outcome for the event "Nausea(Nausea)" was Not recovered.

The outcome for the event "Vomiting(Vomiting)" was Not recovered.

The outcome for the event "A loss of appetite(Appetite lost)" was Not recovered.

The outcome for the event "disgusted by food(Food disgust)" was Not recovered.

The outcome for the event "Ozempic 0.5 mg applied in 18 clicks(Wrong technique in product usage process)" was Not recovered.

Reporter's causality (Ozempic 1.0 mg) -

thyroid was very out of balance (Hypothyroidism aggravated)(Hypothyroidism) : Possible

thyroid was very out of balance (Hypothyroidism aggravated)(Condition aggravated) : Possible

Gastritis (pain in the upper part of the stomach), bad-tasting belching,feel unwell(Gastritis) : Unknown

Nausea(Nausea) : Unknown

Vomiting(Vomiting) : Unknown

A loss of appetite(Appetite lost) : Unknown

disgusted by food(Food disgust) : Unknown

Ozempic 0.5 mg applied in 18 clicks(Wrong technique in product usage process) : Unknown

Company's causality (Ozempic 1.0 mg) -

thyroid was very out of balance (Hypothyroidism aggravated)(Hypothyroidism) : Unlikely

thyroid was very out of balance (Hypothyroidism aggravated)(Condition aggravated) : Unlikely

Gastritis (pain in the upper part of the stomach), bad-tasting belching,feel unwell(Gastritis) : Possible

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

Nausea(Nausea) : Possible
 Vomiting(Vomiting) : Possible
 A loss of appetite(Appetite lost) : Possible
 disgusted by food(Food disgust) : Unlikely
 Ozempic 0.5 mg applied in 18 clicks(Wrong technique in product usage process) : Possible

Reporter Comment: Patient suspended the suspect product as not losing weight and restarted the product on 07-JUN-2025 or 08-JUN-2025

14-19. SUSPECT DRUG(S) continued

14. SUSPECT DRUG(S) (include generic name)	15. DAILY DOSE(S); 16. ROUTE(S) OF ADMIN	17. INDICATION(S) FOR USE	18. THERAPY DATES (from/to); 19. THERAPY DURATION
#1) Semaglutide B 1.34 mg/ml PDS290 1.0 mg (SEMAGLUTIDE 1.34 mg/mL) Solution for injection, 1 mg {Lot # PP5L760; Exp.Dt. JAN-2027}; Regimen #1	18 clicks to get 0.5 mg; Subcutaneous	type 2 diabetes mellitus (Type 2 diabetes mellitus) Obesity (Obesity)	26-MAR-2025 / Unknown; Unknown
#1) Semaglutide B 1.34 mg/ml PDS290 1.0 mg (SEMAGLUTIDE 1.34 mg/mL) Solution for injection, 1 mg {Lot # PP5L760; Exp.Dt. JAN-2027}; Regimen #2	UNK (Resumed); Subcutaneous	type 2 diabetes mellitus (Type 2 diabetes mellitus) Obesity (Obesity)	JUN-2025 / Ongoing; Unknown

22. CONCOMITANT DRUG(S) AND DATES OF ADMINISTRATION continued

#4) XIGDUO (DAPAGLIFLOZIN PROPANEDIOL MONOHYDRATE, METFORMIN HYDROCHLORIDE) Tablet ; MAY-2025 / Ongoing

#6) Novolin R Penfill (INSULIN HUMAN 100 IU/mL) Solution for injection, 100 IU/mL; 2010 / Ongoing

#7) Novolin N (INSULIN ISOPHANE) Suspension for injection ; 2010 / Ongoing

#8) BIOSARTAN (LOSARTAN POTASSIUM) ; 2015 / Ongoing

23. OTHER RELEVANT HISTORY continued

From/To Dates	Type of History / Notes	Description
Unknown to Ongoing	Current Condition	Hypothyroidism (Hypothyroidism);
Unknown to Ongoing	Current Condition	Depression (Depression);
Unknown to Ongoing	Current Condition	Anxiety (Anxiety);
Unknown to Ongoing	Current Condition	Blood pressure high (Hypertension);