

SUSPECT ADVERSE REACTION REPORT

I. REACTION INFORMATION

1. PATIENT INITIALS (first, last) PRIVACY	1a. COUNTRY COSTA RICA	2. DATE OF BIRTH			2a. AGE 70 Years	3. SEX Female	3a. WEIGHT Unk	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	
			PRIVACY					04	NOV	2024	

7 + 13 DESCRIBE REACTION(S) (including relevant tests/lab data)
Event Verbatim [PREFERRED TERM] (Related symptoms if any separated by commas)
Other Serious Criteria: Med sig
Plateletopenia/ Thrombocytopenia [Thrombocytopenia]
Low platelets [Platelet count decreased]
Patient administered Verzenio tablet once daily; No AE [Off label use]

Case Description: This solicited case, reported by a consumer via a Patient Support Program (PSP), with additional information from a second consumer via PSP, concerned a 70-year-old female patient of an unknown origin.

(Continued on Additional Information Page)

II. SUSPECT DRUG(S) INFORMATION

14. SUSPECT DRUG(S) (include generic name) #1) Abemaciclib (Abemaciclib) Tablet [Lot # D669613; Exp.Dt. APR-2026] (Continued on Additional Information Page)		20. DID REACTION ABATE AFTER STOPPING DRUG? <input type="checkbox"/> YES <input checked="" type="checkbox"/> NO <input type="checkbox"/> NA
15. DAILY DOSE(S) #1) 150 mg, bid	16. ROUTE(S) OF ADMINISTRATION #1) Oral	
17. INDICATION(S) FOR USE #1) Breast cancer (Breast cancer)		21. DID REACTION REAPPEAR AFTER REINTRODUCTION? <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> NA
18. THERAPY DATES(from/to) #1) 08-OCT-2024 / 04-NOV-2024	19. THERAPY DURATION #1) 28 days	

III. CONCOMITANT DRUG(S) AND HISTORY

22. CONCOMITANT DRUG(S) AND DATES OF ADMINISTRATION (exclude those used to treat reaction) #1) ARIMIDEX (ANASTROZOLE) Unknown ; Unknown #2) LOPERAMIDE HYDROCHLORIDE (LOPERAMIDE HYDROCHLORIDE) #3) METFORMIN (METFORMIN) Tablet, 500 mg; 2022 / Ongoing #4) IRBESARTAN (IRBESARTAN) Unknown ; 2020 / Ongoing #5) VITAMIN D [COLECALCIFEROL] (COLECALCIFEROL) Unknown ; Unknown #6) CALCIUM (CALCIUM) Unknown ; Unknown (Continued on Additional Information Page)		
23. OTHER RELEVANT HISTORY. (e.g. diagnostics, allergies, pregnancy with last month of period, etc.)		
From/To Dates 2022 to Ongoing Unknown to Ongoing	Type of History / Notes Medical Condition Medical Condition	Description Type 2 diabetes mellitus (Type 2 diabetes mellitus) Arterial hypertension (Hypertension)

IV. MANUFACTURER INFORMATION

24a. NAME AND ADDRESS OF MANUFACTURER Eli Lilly Interamerica Inc (AR Branch) Tronador 4890 - Piso 12 Buenos Aires, Capital Federal CP: 1430 ARGENTINA Phone: 54 1145464000		26. REMARKS
	24b. MFR CONTROL NO. CR202410007986	25b. NAME AND ADDRESS OF REPORTER NAME AND ADDRESS WITHHELD. NAME AND ADDRESS WITHHELD. NAME AND ADDRESS WITHHELD.
24c. DATE RECEIVED BY MANUFACTURER 17-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 23-JUN-2025	25a. REPORT TYPE <input type="checkbox"/> INITIAL <input checked="" type="checkbox"/> FOLLOWUP: 2	

ADDITIONAL INFORMATION

7+13. DESCRIBE REACTION(S) continued

Medical history included diabetes/ type two diabetes since 2022, arterial hypertension, high blood pressure, and low platelet problems since approximately 2022. Concomitant medications included loperamide hydrochloride for the prevention of diarrhea, metformin for diabetes, colecalciferol, olanzapine, amlodipine and calcium for an unknown indication, and irbesartan for high blood pressure.

The patient was prescribed abemaciclib (Verzenio) tablet, 150 mg, twice daily, orally, beginning on 08-Oct-2024, with anastrozole once a day in morning, as concomitant chemotherapy, both for breast cancer. On an unknown date, from the second week of starting abemaciclib therapy, she would start taking 150mg, once daily (off label use). Also, on 04-Nov-2024, she experienced low platelets (values, units and reference ranges were not provided). Due to which, abemaciclib was suspended on 04-Nov-2024. On the week of 20-Dec-2024, abemaciclib therapy was suspended because her platelets were lowered. On 24-Dec-2024, she underwent a test to verify her platelets (no date, results, reference range nor units provided). She did not take any corrective treatment for low platelets. In Mar-2025, she resumed abemaciclib therapy 150 mg every 12 hours. On 15-Mar-2025, she had plateletopenia/ thrombocytopenia. The event of thrombocytopenia was considered as serious by the reporter due to its medical significance reason. On 20-May-2025, a blood test was performed, which showed slightly low platelets (no date, results, reference range and units not provided). Therefore, the doctor decided to discontinue the treatment on 20-May-2025, and instructed her to resume it in 8 days. Outcome of low platelets and thrombocytopenia was not recovered while unknown for other event. Information regarding corrective treatment was not provided. Abemaciclib therapy was ongoing.

The initial reporting consumer did not provide relatedness of the events with abemaciclib therapy.

Update 29-Oct-2024: Information was received from initial reporting consumer on 08-Oct-2024 via PSP. No new medically significant information was received. No changes were made to the case.

Update 24-Nov-2024: Information was received from initial reporting consumer on 13-Nov-2024 and 18-Nov-2024 via PSP. Both follow up processed together. Added one non serious event of platelet count decreased, three concomitant drug anastrozole, vitamin d, calcium, stop date of suspect drug. Updated action taken from unknown to drug discontinued. Updated narrative with new information.

Update 06-Jan-2025: Additional information was received on 27-Dec-2024 from second reporting consumer via a PSP. Added one lab test dated 24-Dec-2024 and a second reporting consumer. Updated age of patient from 56 to 70 years in narrative, platelet count decreased outcome from unknown to not recovered, anastrozole indication from unknown to breast cancer, low platelets corrective treatment from unknown to no, and narrative accordingly.

Update 01-Apr-2025: Additional information received on 27-Mar-2025 from second reporter via PSP. This case was upgraded to serious due to addition of a serious event thrombocytopenia. Added concomitant drugs as olanzapine and amlodipine. Updated event coding for previously captured Intercepted product prescribing error to off label use as patient was prescribed by physician to take 150mg daily . Accordingly updated the narrative with the new information.

Update 26-May-2025: Additional information received on 21-May-2025 and 22-May-2025 from second reporter via PSP. Added one new dosage regimen for abemaciclib therapy and abemaciclib therapy stop date. Accordingly updated the narrative with the new information.

Update 20-Jun-2025: Additional information was received from second reporter via PSP on 17-Jun-2025. Added start date of abemaciclib dosing regimen of 150 mg twice daily. Updated narrative accordingly.

13. Lab Data

#	Date	Test / Assessment / Notes	Results	Normal High / Low
1		Platelet count	low values, units and reference ranges were not provided.	
2	24-DEC-2024	Platelet count	(no date, results, reference range nor units provided)	

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
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ADDITIONAL INFORMATION

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) Abemaciclib (Abemaciclib) Tablet; Regimen #2	150 mg, daily; Oral	Breast cancer (Breast cancer)	Unknown; Unknown
#1) Abemaciclib (Abemaciclib) Tablet; Regimen #3	150 mg, daily; Oral	Breast cancer (Breast cancer)	Unknown; Unknown
#1) Abemaciclib (Abemaciclib) Tablet; Regimen #4	150 mg, bid; Oral	Breast cancer (Breast cancer)	MAR-2025 / 20-MAY-2025; Unknown

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

#2) LOPERAMIDE HYDROCHLORIDE (LOPERAMIDE HYDROCHLORIDE) Unknown ; 08-OCT-2024 / Ongoing

#7) TELORZAN (OLANZAPINE) Unknown ; Unknown

#8) AMLODIPINE (AMLODIPINE) Unknown ; Unknown

23. OTHER RELEVANT HISTORY continued

From/To Dates	Type of History / Notes	Description
Unknown to Ongoing	Medical Condition	Blood pressure high (Hypertension);
2022 to Unknown	Medical Condition since 1 or 2 years	Low platelets (Platelet count decreased);