Humana

Pharmacy Solutions®

Drug recall notice for Steqeyma (ustekinumab-stba) 90 mg/mL

To assist you in the care of your patients, Humana is alerting you to the recall of Steqeyma® (ustekinumab-stba) 90 mg/mL on Aug. 13, 2025.[1] We recommend you review your medical records and contact all patients for whom you have prescribed this medication to warn them of the recall.

The drug manufacturer, Celltrion USA, is voluntarily recalling one lot of Steqeyma 90 mg/mL to the consumer level as a precautionary measure due to an incorrect assembly issue identified during quality control.

To date, Celltrion USA has received no reports of adverse events related to this recall.

Medications included in this recall
Visit the recall website for specific details about the recalled medication.

Product name	National Drug Code	Lot number	Expiration date
Steqeyma 90 mg/mL	72606-0028-01	4N2P07S51	Nov. 30, 2027

Information for providers:[1]

- We sent a letter to your Humana-covered patients with a claim for Steqeyma 90 mg/mL and asked them to contact their providers if their medication is included in the recall and if they have experienced problems that could be related to using these drug products.
- Providers with questions can contact:
  - Celltrion USA by phone at 201-241-0038, Monday – Friday, 9 a.m. – 5 p.m., Eastern time or by email at Batsheva.bain@celltrionhc.com.
- Patients can report adverse reactions or quality problems experienced with the use of these products to the FDA's MedWatch Adverse Event Reporting Program online, by phone or by fax.
  - **Online:** Submit the report.
    - Select "Form FDA 3500 - Voluntary Reporting."
  - **Phone or fax:** Download the form.
    - Complete and submit "Form FDA 3500 - Voluntary Reporting " by phone at 800-FDA-1088 (332-1088) or by fax to 800-FDA-0178 (332-0178).

Reference

1. "Steqeyma 90 mg/mL," California State Board of Pharmacy, last accessed Oct. 13, 2025, https://www.pharmacy.ca.gov/about/recall_alerts/081325_celltrion.pdf.