Humana.

Drug recall for Steqeyma® (Ustekinumab-stba) 90 mg/mL injection

On August 13, 2025, Celltrion USA, initiated a voluntary recall of one (1) lot of Steqeyma® (Ustekinumab-stba) 90 mg/mL injection to the consumer level, due to incorrect assembly.

Please carefully review the impacted medication and talk to your doctor about it right away.

What this means to you:

- To date, the pharmaceutical company has not received any adverse event reports related to this recall.
- Talk to your doctor or healthcare provider about switching to another medication or obtaining the same medication that is not part of the recall.
- To determine if your medication is affected, you should look at the drug name and company name on the label of your prescription. If the information is not on the package, you should contact the pharmacy that dispensed the medication.
- Please refer to the recall notice for the most current updates to this drug recall.
- You can also contact Celltrion USA at 1-201-241-0038, Monday to Friday from 9:00 am to 5:00 pm ET.
- You should contact your doctor or healthcare provider if you have experienced any problems that may be related to taking or using these drug products.