

## **Drug Recall**

**DATE OF RECALL:** November 22, 2021

**DRUG NAME:** Levetiracetam Injection 500mg/5 mL (100mg/1 mL)

**RECALLING FIRM:** Sagent Pharmaceuticals, Inc.

**REASON FOR RECALL:** This recall was issued due to a container closure integrity issue found in reserve sample vials that may result in a non-sterile product

**FDA LINK:** <a href="https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/sagent-pharmaceuticals-inc-issues-voluntary-nationwide-recall-levetiracetam-injection-usp-due-lack">https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/sagent-pharmaceuticals-inc-issues-voluntary-nationwide-recall-levetiracetam-injection-usp-due-lack</a>

**OTHER DETAILS:** N/A

## **RECALLED PRODUCT:**

Product Description	NDC Number	Affected Lots	<b>Expiration Date</b>
Levetiracetam Injection	25021-0780-05	B0G85VB	06/2022
500mg/5 mL (100mg/1 mL)			
Levetiracetam Injection	25021-0780-05	B0K88VA	09/2022
500mg/5 mL (100mg/1 mL)			
Levetiracetam Injection	25021-0780-05	B0K89VA	09/2022
500mg/5 mL (100mg/1 mL)			
Levetiracetam Injection	25021-0780-05	B1G194A	06/2023
500mg/5 mL (100mg/1 mL)			