

## **Drug Recall**

DATE OF RECALL: February 3, 2025

**DRUG NAME:** fentanyl transdermal system

**RECALLING FIRM:** Alvogen

**REASON FOR RECALL:** Alvogen announced a voluntary, consumer level recall of one lot of fentanyl transdermal system 25 mcg/h because there is a potential that patches could be multi-stacked, adhered one on top of the other, in a single product pouch.

**FDA LINK:** <a href="https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/alvogen-issues-voluntary-nationwide-recall-one-lot-fentanyl-transdermal-system-25-mcgh-due-defective">https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/alvogen-issues-voluntary-nationwide-recall-one-lot-fentanyl-transdermal-system-25-mcgh-due-defective</a>

**OTHER DETAILS:** N/A

## **RECALLED PRODUCT:**

Product Description	NDC Number	Lot Number	Expiration Date
Fentanyl Transdermal System, 25 mcg/h; Five individually wrapped and labeled pouches	47781-424-47	108319	4/2027