The Board of Pharmacy has received notice of the following product recall. The Board strongly encourages pharmacies to immediately review their quality assurance and recall policies and procedures to determine if any corrective action is required.

Alvogen, Inc. is voluntarily recalling one lot of Fentanyl Transdermal System 25 mcg/h transdermal patches to the consumer level. A small number of patches is impacted. The reason for the recall is that there is a potential that patches could potentially be multi-stacked, adhered one on top of the other, in a single product pouch. This transdermal system is manufactured by Kindeva Drug Delivery L.P., Northridge, CA and is distributed by Alvogen, Inc. as a private label distributor.

Shipments of the affected lot were sent between 06/11/2024 and 10/21/2024. There is a possibility that the application of a multi-stacked 25 mcg/h patch could result in serious, life
threatening, or fatal respiratory depression. Groups at potential increased risk could include first-time
recipients of such patches, children, and the elderly.

PRODUCTS: Fentanyl Transdermal System 25mcg/hr
Pack Sizes: Carton (5 pouches/carton) Pouch (1 patch/pouch)

NDC NUMBERS: Carton: 47781-424-47
Pack: 47781-424-11

LOT NUMBER: 108319

EXPIRATION DATE: 04/2027

SHIP DATES: 06/11/24 – 10/21/24