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.

This recall has been initiated by Sun Pharmaceutical Industries, Inc. in response to cGMP deviations. Lot DND1515A is being recalled out of an abundance of caution. Per FDA recommendation dated June *5,* 2023, this recall is revised from the depth of wholesale level to retail level. Sun Pharmaceutical Industries, Inc. initiated shipment of this product on Dec 14, 2022.

Product: Buprenorphine Hydrochloride Sublingual Tablets. 8 mg, 30 count

Lot Number: DND1515A

NDC Number: 62756-460-83

Expiration Date: 08/2024