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.

Amneal Pharmaceuticals LLC is recalling 20 lots of Dexmethylphenidate Hydrochloride Extended-Release Capsules, 10 mg (NDC 0115-9919-01) 100 count, due to the presence of FD&C Red No. 2 (amaranth), a color additive that is no longer authorized for use by the U.S. Food and Drug Administration (FDA). No other strengths are impacted on this recall.

Dexmethylphenidate HCL ER Capsules is indicated for the treatment of Attention Deficit Hyperactivity Disorder (ADHD).

The health risk associated with ingestion of low levels of FD&C Red No. 2 (amaranth) depends on the amount consumed, frequency of exposure, and individual susceptibility. Animal studies have identified adverse effects, including kidney changes, following repeated exposure at higher doses; however, these findings do not establish that low-dose ingestion causes similar harm in humans.

PRODUCT: Dexmethylphenidate Hydrochloride Extended-Release Capsules, 10 mg

NDC NUMBER: 0115-9919-01

Lot Number	Expiration Date	First Ship Date
BA12225A	11/2026	3/4/2025
BA12325A	11/2026	3/4/2025
BA12425A	11/2026	2/20/2025
BA12525A	11/2026	2/13/2025
BA36225A	01/2027	3/19/2025
BA36325A	01/2027	3/12/2025
BD22925A	03/2027	7/15/2025
BD23025A	03/2027	7/1/2025
BD23125A	03/2027	7/1/2025

BD23225A	03/2027	7/8/2025
BD23325A	04/2027	7/17/2025
BE03025A	04/2027	7/23/2025
BE03125A	04/2027	7/30/2025
BE03225A	04/2027	7/17/2025
BK14325A	10/2027	1/5/2026
BK14425A	10/2027	1/5/2026
BK14525A	10/2027	1/21/2026
BK14625A	10/2027	1/15/2026
BA13326A	01/2028	3/26/2026
BA13426A	01/2028	3/30/2026