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.

NDC	Lot	Exp. Date	Bottle Size
00555-0808-02	100022971	08/2022	100 Capsules
00555-0808-02	100022972	05/2023	100 Capsules
00555-0808-02	100028920	09/2023	100 Capsules

Tretinoin Capsules 10 mg

Teva Pharmaceuticals USA Inc. is initiating a nationwide recall of three lots of Tretinoin Capsules 10 mg to the **Retail** Level in the United States. The affected product lots were distributed under the label of Teva Pharmaceuticals USA Inc. Teva USA distribution records indicate the above referenced lots shipped from 09/01/2021 through 03/07/2022.

This recall has been initiated because the dissolution results for Lot 100028920 are below specification limit. The remaining two lots in this recall may potentially be implicated with this problem since they are manufactured and tested in the same way and with the same materials. It should be noted that all test results were within specification limits, including dissolution, at the time of product release for all of the lots in this recall.