The Board of Pharmacy has received notice of the following product recall:

Strength	NDC	Pack Size	Description
20 mg / 12.5 mg	68180-519-02	500's bottle	Lisinopril and Hydrochlorothiazide Tablets USP, 20 mg/12.5 mg are yellow, round tablets, with "LL" debossed on one side and "B02" on other side.

Lupin Pharmaceuticals Inc. is recalling one lot - **H801815, expiry March 2021 -** of Lisinopril and Hydrochlorothiazide Tablets USP, 20mg/12.5mg to the retail level. As an abundance of caution, this product lot is being recalled due to a product complaint where one tablet of Lupin's Fenofibrate 145mg was observed in the 500's count product bottle. Any potential health hazard is very unlikely. The recalled lot was distributed between August 23, 2018, and August 30, 2018, to wholesalers, distributors, drug chains, mail order pharmacies and super markets nationwide.

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