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	Product Name, Strength and Package Count	Batch Number	Expiration Date
13668-159-30	Telmisartan and Hydrochlorothiazide Tablets, USP 40/ 12.5 mg, 30 Count	BZ74G001	12/2021

Torrent Pharmaceuticals Limited is initiating a recall of Telmisartan and Hydrochlorothiazide Tablets USP, 40/ 12.5 mg to the Retail level. This recall is based upon a high out of specification Assay result of Hydrochlorothiazide detected during routine stability testing for this batch. Considering all available information, it is determined that the slight increase in Assay would not have a clinically significant impact on safety and efficacy. There were no adverse event or product complaint cases received for this batch. Hence, based on available data, there is a minimal chance of health hazard.