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.

Dr. Reddy's Laboratories Inc. is issuing a recall as a precautionary measure due to OOS results observed at 24-month long term stability condition (25°C/60% RH) in Dissolution test (sample did not meet S3 stage criteria for dissolution). Risk Assessment: The lower dissolution results may affect the pharmacokinetic profile i.e., the desired therapeutic concentration of Tizanidine may be low. Therefore, the probability of lack of desired effect may be observed in a patient who may consume Tizanidine Tablets. The product Distribution Dates: May 24, 2021 – August 3, 2021. Refer to the product lot to be recalled in the table below.

Product Description	**Lot Number**	**Expiry Date**	**NDC Number**
Tizanidine Tablets USP, 4mg, 1000 Ct	T2100585	03/2024	55111-180-10
Tizanidine Tablets USP, 4mg, 1000 Ct	T2100586	03/2024	55111-180-10
Tizanidine Tablets USP, 4mg, 1000 Ct	T2100587	03/2024	55111-180-10