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.

This recall is being re-issued following reclassification from Class Ill to Class II by the U.S. Food and Drug Administration. Please notify your customers accordingly.

This is to inform you of a product recall involving Metoprolol Succinate Extended-Release (ER) Tablets USP, 25 mg by Ascend Laboratories.

An out-of-specification (OOS) result was observed during dissolution test analysis during long term stability testing (LT conditions: 25 ± 2 •c / 60 ± 5 % RH) of Metoprolol Succinate ER Tablets USP, 25 mg, for batch number 25140859.

Metoprolol Succinate is indicated for hypertension in patients six years of age and older as well as for angina pectoris and heart failure in adult patients. The dissolution data show a limited acceleration of drug release confined to the 8-hour time point, while all other intervals and 12-month stability results remain within specification, confirming the integrity of the extended-release profile. As this represents a marginal terminal-phase release acceleration without evidence of dose dumping or uncontrolled delivery, the deviation is not expected to be clinically significant and constitutes a minor health hazard at most.

Our firm began shipping this product on June 17, 2025.

PRODUCT: Metoprolol Succinate Extended-Release (ER) Tablets USP, 25 mg, 100 tablets per bottle

NDC NUMBER: 67877-590-01

LOT NUMBER: 25140859

INITIAL DISTRIBUTION DATE: 6/17/2025

EXPIRY DATE: 01/31/2027