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.

CLARIFICATION OF AFFECTED PRODUCT

Description	Lot # and Exp Date	NDC	UPC
ZONISAMIDE CAP 100MG GLEN 100@	29200049 08-31-23; 29200015 03-31-23; 29200016 03-31-23; 29200030 05-31-23; 29200031 05-31-23; 29200032 05-31-23; 29200033 06-30-23; 29200037 06-30-23; 29200038 06-30-23; 29200039 07-31-23; 29200041 07-31-23; 29200014 02-28-23; 29200048 08-31-23; 29200050 08-31-23; 29200072 11-30-23; 29200073 11-30-23; 29200074 11-30-23; 29200075 11-30-23; 29200076 11-30-23; 29200042 07-31-23	68462013001	36846213001
ZONISAMIDE CAP 100MG GLEN 500@	29200014 02-28-23; 29200015 03-31-23; 29200016 03-31-23	68462013005	36846213005

Description	Lot # and Exp Date	NDC	UPC
ZONISAMIDE CAP 50MG GLEN 100@	29190045 05-31-22; 29190043 05-31-22; 29190044 05-31-22	68462012901	36846212901

Note: Updated recall notification from Glenmark does not include NDC 68462012801 as part of this recall; although the NDC was listed in the original notice, no affected lots were listed.

Glenmark is voluntarily recalling the above items/lots due to gaps in the quality system in the Quality Control microbiology laboratory. This recall is to the retal level. Affected product started shipping September 24, 2019.