Zonisamide - Glenmark

Glenmark announced a consumer-level recall of several lots of Zonisamide capsules due to gaps in the microbiology quality control system or due to lack of stability data.

Product	Zonisamide
Manufacturer	Glenmark
Recall identification date	04/28/2022
Affected NDCs	68462-130-01, 68462-130-05, 68462-128-01, 68462-129-01

Consumers should contact their physician or healthcare provider if they have experienced any problems that may be related to taking or using this product.