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	Expiration Date	Pack Size	Lot#	Distribution Dates
67877-545-68	Cephalexin FOS USP	250mg/5ml	March-2021	200ml	19141869	7/9/2019 to 7/18/2019
67877-545-68	Cephalexin FOS USP	250mg/5ml	March-2021	200ml	19141870	7/3/2019 to 9/9/2019
67877-545-68	Cephalexin FOS USP	250mg/5ml	May-2021	200ml	19142762	11/3/2019 to 12/27/2019
67877-545-68	Cephalexin FOS USP	250mg/5ml	July-2021	200ml	19143826	2/6/2020 to 2/20/2020
67877-545-68	Cephalexin FOS USP	250mg/5ml	July-2021	200ml	19143923	1/24/2020 to 2/7/2020
67877-545-68	Cephalexin FOS USP	250mg/5ml	July-2021	200ml	19143941	1/2/2020 to 2/7/2020
67877-545-68	Cephalexin FOS USP	250mg/5ml	July-2021	200ml	19143954	11/18/2019 to 11/26/2019
67877-545-88	Cephalexin FOS USP	250mg/5ml	September-2021	100ml	19144841	3/10/2020 to 3/24/2020
67877-545-88	Cephalexin FOS USP	250mg/5ml	April -2022	100ml	20141673	9/28/2020 to 10/2/2020

Ascend Laboratories has initiated this recall as a precautionary measure. During related substance test analysis of Cephalexin for oral suspension, USP 250mg/5ml, results of Individual Unidentified impurity are found on higher side but within the specification limit. This recall should be carried out to the Retail level.