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.

Alpha$_1$proteinase inhibitor (Human)

ZEMAIRA® - 1000 mg Range

NDC Number: 0053-7201-02

Batch	Expiry Date		Batch	Expiry Date		Batch	Expiry Date
Y411604	3/31/2021		P100053388	1/29/2022		P100128011	8/18/2022
Y412908	7/31/2021		P100063362	2/14/2022		P100134128	9/8/2022
Y413309	9/7/2021		P100066936	2/17/2022		P100134132	9/12/2022
Y413509	9/13/2021		P100066944	2/22/2022		P100169473	12/11/2022
Y413609	9/15/2021		P100074924	3/31/2022		P100171844	12/15/2022
Y414009	9/26/2021		P100125236	3/31/2022		P100171848	1/6/2023
Y414210	10/5/2021		P100125239	4/22/2022		P100171852	1/24/2023
Y414310	10/9/2021		P100089996	4/22/2022		P100175268	1/25/2023
Y414410	10/15/2021		P100125241	4/28/2022		P100246358	7/17/2023
Y415112	12/9/2021		P100125242	5/14/2022		P100246366	7/21/2023
Y415412	12/19/2021		P100122777	6/1/2022		P100250559	7/26/2023
P100049736	1/13/2022		P100107523	6/19/2022		P100248400	7/29/2023
P100051365	1/20/2022		P100124006	8/13/2022		P100250563	8/6/2023

CSL Behring has instituted a recall of ZEMAIRA® at the hospital/pharmacy level, due to a manufacturing deviation that occurred during the filling process.