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 is to inform you that Grifols Therapeutics is initiating a withdrawal of two lots of Gamunex® -C 10%, 20G Vial, as detailed below.

Grifols is committed to providing the highest quality medicines in the market. This withdrawal is being conducted as a precautionary measure due to an increased rate of allergic/hypersensitivity type reactions associated with these specific lots. A small number of the reactions were considered medically significant. Hypersensitivity and anaphylactic/anaphylactoid reactions are a known risk with immune globulin products.

This withdrawal is being conducted with the knowledge of the U.S. Food and Drug Administration, Center for Biologics Evaluation and Research. This withdrawal is required to be conducted to the consumer/user level.

The Gamunex® -C 10% lots affected by this withdrawal are:

Lot Number	Material Number	Expiration Date	Market Release	NDC Number
B01J093362	729688	02 DEC 2027	14 FEB 2025	13533-800-24
B01J094362	729688	08 DEC 2027	13 FEB 2025	13533-800-24