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.

Eli Lilly and Company is recalling to the patient (consumer/user) level one specific lot of Glucagon Emergency Kit in cooperation with the US Food and Drug Administration. The recall extends to lot **D239382D.**

Product Description	**NDC Number**	**Lot # / Exp Date**
MS8031001AM Glucagon Emergency Kit for Low Blood Sugar Glucagon for jection, 1 mg per vial Diluent for Glucagon, 1 mL syringe	NDC 0002-8031-01	D239382D/ 04 2022

Eli Lilly and Company is recalling lot D239382D, Expiration April 2022, of Glucagon Emergency Kit for Low Blood Sugar (Glucagon for Injection, 1 mg per vial: Diluent for Glucagon, 1 mL syringe), to the consumer/user level. Lilly is recalling lot D239382D to the patient level because of a product complaint reporting that the vial of Glucagon was in liquid form instead of the powder form. The firm' s investigation indicates that the liquid in this Glucagon vial could be related to the manufacturing process. The use of the liquid form of this product may fail to treat severe low blood sugar due to loss of potency.

The Eli Lilly distribution period of this lot D239382D is from 29 December 2020 to 25 March 2021.