# Glucagon Emergency Kit

This notice is to inform you that on September 26th, 2021 the U.S. Food and Drug Administration (FDA) announced that Eli Lilly and Company is voluntarily recalling one lot of Glucagon Emergency Kit for Low Blood Sugar due to a product complaint reporting that the vial of Glucagon was in liquid form instead of the powder form. The use of the liquid form of this product may fail to treat severe low blood sugar due to loss of potency.

Risk Statement: Severe hypoglycemia in patients with diabetes, if not reversed, can potentially cause adverse health consequences ranging from transient, minor complaints to neurological damage, seizures, and even death if not promptly treated. Associated with the one product complaint, it was reported to Lilly that the involved patient experienced lack of drug effect and also reported subsequent seizures.

Glucagon Emergency Kit is used as an anti-hypoglycemic agent and a gastrointestinal motility inhibitor indicated for the treatment of severe hypoglycemia in pediatric and adult patients with diabetes mellitus.

Consumers in possession of Glucagon Emergency Kit lot D239382D should contact The Lilly Answers Center at 1-800-LILLYRX (1-800-545-5979) for return and replacement instructions for the product (hours of operation are Monday- Friday, 9AM – 7PM EST) and should contact their health care provider for guidance. Consumers should contact their physician or healthcare provider if they have experienced any problems that may be related to taking or using this product.