Mass General Brigham
Health Plan

Drug Recall

DATE OF RECALL: November 9, 2022

DRUG NAME: Octreotide Acetate Injection 500mcg/ml (Syringe)

RECALLING FIRM: Mylan Institutional LLC

REASON FOR RECALL: This recall was issued due to a product complaint for the presence of glass particles in a syringe.

FDA LINK: Mylan Institutional LLC, a Viatris Company, Issues a Voluntary Recall of One Lot of Octreotide Acetate Injection, 500 mcg/mL, Due to Glass Particulates in a Syringe | FDA

OTHER DETAILS: See FDA Link

RECALLED PRODUCT:

Affected Product	NDC #	Lot Number
Octreotide Acetate Injection 500mcg/ml (Syringe), carton	67457-0246-01	AJ21002 exp. 03/2024
Octreotide Acetate Injection 500mcg/ml (Syringe), single	67457-0246-00	