Mass General Brigham
Health Plan

Drug Recall

DATE OF RECALL: April 25, 2023

DRUG NAME: FYREMADEL® (ganirelix acetate) Injection, 250 mcg/0.5 mL

RECALLING FIRM: Sun Pharmaceutical Industries Ltd

REASON FOR RECALL: This recall was issued because of a product quality complaint stating, “glass particle observed in prefilled syringe of FYREMADEL® (ganirelix acetate) Injection 250 mcg/0.5 mL, lot number HAD1190A”.

FDA LINK: N/A

OTHER DETAILS: N/A

RECALLED PRODUCT:

Product Description	NDC Number	Lot Number	Expiration Date
FYREMADEL® (ganirelix acetate) Injection, 250 mcg/0.5 mL	55566-1010-01	HAD1190A	02/2024