Mass General Brigham
Health Plan

Drug Recall

DATE OF RECALL: August 11, 2026

DRUG NAME: Tyenne Injection

RECALLING FIRM: Fresenius Kabi

REASON FOR RECALL: Fresenius Kabi announced a voluntary, consumer level recall of one lot of Tyenne (tocilizumab-aazg) injection 400 mg/20 mL (20 mg/mL) because the product was found to contain glass particles.

FDA LINK: Fresenius Kabi Issues Nationwide Recall of Tyenne® (tocilizumab-aazg) Injection 400 mg/20 mL Vials Due to Presence of Glass Particles | FDA

OTHER DETAILS: N/A

RECALLED PRODUCT:

Product Description	NDC Number	Lot Number	Expiration Date
Tyenne (tocilizumab-aazg) Injection, 400 mg/20 mL(20mg/mL), 20 mL vial	65219-594-20	16UI07	08/2028