California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

Integra LifeSciences Corp. AURORA Surgiscope System

Recall Date	Product Description	Recalling Firm	Recall Reason
03/06/2025	AURORA Surgiscope System, Sterile, single use device that contains a Sheath, Obturator, and Imager. Manufacturer's Product Numbers (Catalog Number): (1) ASX15/60 and (2) ASX15/80	Integra LifeSciences Corp.	Possibility for the obturator to break (separate).

Recall Class	Product Identification	Distribution	Affected Dates
II	Catalog Number: (1) ASX15/60 (2) ASX15/80 UDI-DI: (1) 00850002332254 (2) 008500023332247 Lot No.: (1) All unexpired lots (2) All unexpired lots	116 units in California	February and prior

For additional information, please visit the FDA Website.