California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

Intersurgical Inc Recalls i-view video laryngoscope

Recall Date	Product Description	Recalling Firm	Recall Reason
7/11/25	i-view video laryngoscope. Model Number: 8008000. The i-view video laryngoscope is indicated to facilitate laryngoscopy.	Intersurgical Inc	Potential for faulty devices as a result of depleted batteries.

Recall Class	Product Identification	Distribution	Affected Dates
I	Model Number: 8008000. UDI-DI (Case): 05030267166497. UDI-DI (Single Unit): 5030267150649. Lot Numbers: 1240555, 1240793.	US distribution AZ, CA, FL, GA, IL, IN, KS, MD, MI, MO, NV, NY, OH, PA, SC, TX, VA.	June 2025 and prior.

For additional information, please visit the FDAWebsite