State of California—Health and Human Services Agency
California Department of Public Health

Food and Drug Branch – Device Recalls

California Device Recall Information Sheet

Covidien Recalls Mcgrath Mac Video Laryngoscope

Recall Date	Product Description	Recalling Firm	Recall Reason
8/28/2024	**Mcgrath Mac Video Laryngoscope** McGrath Mac Video Laryngoscope, REF: 300-000-000, Non-Sterile, Rx Only	**COVIDIEN** Boulder, CO	Battery management system within Laryngoscope devices may deplete below the deign threshold which may result in thermal event followed by risk of explosion.

Recall Class	Product Identification	Distribution	Affected Dates
I	**Mcgrath Mac Video Laryngoscope** UDI: 15060272980020 05060272980023 F0602729800201	75,544 units Units Nationwide including C	August 2024 and Prior

For additional information, please visit the FDA website.

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