State of California—Health and Human Services Agency

California Department of Public Health

Food and Drug Branch – Device Recalls

CALIFORNIA DEVICE RECALL INFORMATION SHEET

GE Healthcare Recalls MRI Software due to Slicing Error

Recall Date	Product Description	Recalling Firm	Recall Reason
2/22/2023	**EVIS EXERA III Colonovideoscope** For endoscopy and endoscopic surgery within the lower digestive tract (including the anus, rectum, sigmoid colon, colon, and ileocecal valve) Model/Serial: CF-Q180AL	**Olympus Corporation of the Americas** Center Valley, Pennsylvania	A single CF-Q180AL colonovideoscope was utilized in a veterinary endoscopy procedure in advance of being assigned to a medical facility as a service loaner in error, potential for microbial contamination

Recall Class	Product Identification	Distribution	Affected Dates
II	**EVIS EXERA III** UDI: 04953170202315 Serial number 2807443	1 unit in California	December 2022 and prior

FOR ADDITIONAL INFORMATION, PLEASE VISIT THE FDA WEBSITE

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