State of California—Health and Human Services Agency

California Department of Public Health

Food and Drug Branch – Device Recalls

CALIFORNIA DEVICE RECALL INFORMATION SHEET

Olympus Corporation Of The Americas Recalls The Air/Water Valve Provided/Used With The Endoscopes For the loss of one-way valve functionality.

Recall Date	Product Description	Recalling Firm	Recall Reason
8/30/2023	**Olympus Corporation Of The Americas** The Air/Water Valve provided/used with the following Endoscopes: ULTRASONIC GASTRO-FIBERSCOPE, ULTRASONIC GASTROVIDEOSCOPE, ULTRASONIC COLONOV-IDEOSCOPE: GF-UC140P-AL5, GF-UCT140-AL5, GF-UE160-AL5, GF-UCT180, GF-UM20, GF-UM130, GF-UMQ130, GF-UM160, GF-UC160P-OL5,GF-UCT160-OL5	**Olympus Corporation Of The Americas** Center Valley, Pennsylvania	The air/water valve MAJ-1444 used with OER-Pro and OER-Elite may become damaged and result in loss of the oneway valve functionality.

Recall Class	Product Identification	Distribution	Affected Dates
II	**Olympus Corporation Of The Americas.** The Air/Water Valve UDI-DI: 04953170355929. All lot numbers	29590 Units Nationwide	July 2023 and prior

FOR ADDITIONAL INFORMATION, PLEASE VISIT THE FDA WEBSITE

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