



California Medical Device Recall Information



Recall Name

Verathon Recalls GlideScope Titanium Single-Use Video Laryngoscope Due to Disruption of Video Feed

Recall Date	Product Description	Recalling Firm	Recall Reason
1/29/16	GlideScope Titanium Single-Use Video Laryngoscope	Verathon, Inc. Bothell, WA	<i>Potential for disruption in the video feed from the camera in the laryngoscope blades to the monitor.</i>
Recall Class	Product Identification	Distribution	Affected Dates
I	Lot Numbers: <ul style="list-style-type: none"> • LoPro S3: 081814 - 093015; • LoPro S4: 081114 - 090315; • MAC S3: 080814 - 101315; • MAC S4: 022514 – 082115 	CA , nationwide	Manufactured: November, 2014 to December, 2015 Distributed: November 14, 2014 to December 29, 2015

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm491325.htm>