



Food and Drug Branch – Device Recalls

**CALIFORNIA DEVICE RECALL INFORMATION SHEET**

**Datascope Recalls Cardiosave Intraaortic Balloon Pump For Autofilling Failure**

Recall Date	Product Description	Recalling Firm	Recall Reason
9/6/2023	<b>Cardiosave Hybrid Intra-Aortic Balloon Pumps (Iabp)</b> Part Numbers 0998-00-0800-31, 0998-UC-0800-31, 0998-00-0800-32, 0998-00-0800-33, 0998-UC-0800-33, 0998-00-0800-34, 0998-00-0800-35, 0998-00-0800-45, 0998-00-0800-52, 0998-UC-0800-52, 0998-00-0800-53, 0998-UC-0800-53, 0998-00-0800-55, 0998-UC-0800-55, 0998-00-0800-65	<b>DATASCOPE CORP.</b> Mahwah, New Jersey	Users were identifying autofill failure conditions on the devices causing pump stops.
9/6/2023	<b>Cardiosave Rescue Intra-Aortic Balloon Pumps (Iabp)</b> Part Numbers 0998-00-0800-75, 0998-00-0800-83, 0998-00-0800-85	<b>DATASCOPE CORP.</b> Mahwah, New Jersey	Users were identifying autofill failure conditions on the devices causing pump stops.

Recall Class	Product Identification	Distribution	Affected Dates
I	<b>Cardiosave Hybrid Intra-Aortic Balloon Pumps (Iabp)</b> All Lot Numbers. Model: 0998-00-0800-31, UDI: 10607567109053; Model: 0998-UC-0800-31, UDI: N/A; Model: 0998-00-0800-32, UDI: 10607567111117; Model: 0998-00-0800-33, UDI: 10607567109008; ...	9175 Units Nationwide	July 2023 and prior

I	<b>Cardiosave Rescue Intra-Aortic Balloon Pumps (Iabp)</b> All Lot Numbers. Model: 0998-00-0800-75, UDI: 10607567112312; Model: 0998-00-0800-83, UDI: 10607567108407; Model: 0998-00-0800-85, UDI: 10607567113449	9175 Units Nationwide	July 2023 and prior
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FOR ADDITIONAL INFORMATION, PLEASE VISIT THE [FDA WEBSITE](#)

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CDPH Food and Drug Branch  
MS 7602 ● P.O. Box 997435 ● Sacramento, CA 95899-7435  
(916) 650-6500 ● (916) 650-6650 FAX  
Internet Address: [www.cdph.ca.gov](http://www.cdph.ca.gov)

