



# California Medical Device Recall Information



## Recall Name

### Datascope Corp / MAQUET Recalls Intra-Aortic Balloon Pumps Due to Potential Mechanical Failure Causing Shut Down

Recall Date	Product Description	Recalling Firm	Recall Reason
03/21/14	Intra-Aortic Balloon Pumps (IABPs) [sold under <b>Datascope Corp. System</b> brand names]	<b>Datascope Corp/ MAQUET</b> Wayne, NJ	<i>Potential mechanical failure of the fan assembly associated with the power supply.</i>
Recall Class	Product Identification	Distribution	Affected Dates
I	<ul style="list-style-type: none"> <li><u>98/98XT Part Numbers:</u> 0998-00-0446-xx 0998-UC-0446-xx 0998-00-0479-xx 0998-UC-0479-xx</li> <li><u>CS100/CS100i Part Numbers:</u> 0998-00-3013-xx 0998-UC-3013-xx 0998-UC-0446Hxx 0998-UC-0479Hxx</li> <li><u>CS300 Part Numbers:</u> 0998-00-3023-xx 0998-UC-3023-xx</li> </ul> <p>[For a list of affected IABP Serial Numbers, contact a MAQUET Service Representative.]</p>	CA, nationwide	Manufactured between:  January 1, 2003 and June 30, 2011

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm396821.htm>