



# California Medical Device Recall Information



## Recall Name

**HeartWare Recalls Batteries used on HeartWare Ventricular Assist Device (HVAD)  
Due to Premature Power Depletion**

Recall Date	Product Description	Recalling Firm	Recall Reason
1/07/16	Batteries: <ul style="list-style-type: none"> <li>used on HeartWare Ventricular Assist Device (HVAD)</li> <li>Model number: <b>1650</b></li> </ul>	<b>HeartWare, Inc.</b> Miami Lakes, FL	<i>The batteries may lose power prematurely due to faulty cells.</i>
Recall Class	Product Identification	Distribution	Affected Dates
I	Serial Numbers:  <b>BAT000001</b> to <b>BAT199999</b>	<b>CA</b> , nationwide	Manufacturing dates: May 19, 2013 to July 1, 2015  Distribution dates: May 21, 2013 to July 31, 2015

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm509185.htm>