



# California Medical Device Recall Information



## Recall Name

**Greatbatch, Inc. Recalls Standard Offset Cup Impactors  
Due to Lack of Sterility Assurance**

Recall Date	Product Description	Recalling Firm	Recall Reason
01/29/14	<b>Standard Offset Cup Impactors</b>	<b>Greatbatch, Inc.</b> Plymouth, MN	<i>The sterilization recommendation in the Instructions For Use did not meet requirements for sterility assurance, which has the potential to result in surgical infection.</i>
Recall Class	Product Identification	Distribution	Affected Dates
I	<u><a href="#">Affected Product List</a></u>	CA, nationwide	Manufactured and distributed from:  2004 to 2013.

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm388883.htm>