



California Medical Device Recall Information



Recall Name

DePuy Orthopaedics Recalls LPS Diaphyseal Sleeve Due to Fracture of the Sleeve

Recall Date	Product Description	Recalling Firm	Recall Reason
2/15/13	The Limb Preservation System (LPS) Diaphyseal Sleeve	DePuy Orthopaedics, Inc. Warsaw, IN	<i>Potential device failure including the fracturing of the Sleeve, leading to loss of limb, infection, or death.</i>
Recall Class	Product Identification	Distribution	Affected Dates
I	LPS Diaphyseal Sleeve Part #: <ul style="list-style-type: none"> • 1987-20-018 • 1987-20-020 • 1987-20-024 • 1987-20-028 All Lots Recalled	CA , nationwide	Manufacturing and distribution dates: From 2008 to July 20, 2012

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm340708.htm>