



California Medical Device Recall Information



Recall Name

Insulet Corporation Recalls OmniPod Insulin Management System Due to Failures That May Result in Inaccurate Dosage

Recall Date	Product Description	Recalling Firm	Recall Reason
7/13/15	OmniPod Insulin Management System Catalog Number: • POD-ZXP420	Insulet Corporation Billerica, MA	<i>Potential failure of the device where the tube fails to fully insert into skin or completely retracts after insertion, and a failure in which the Pod will not pump insulin.</i>
Recall Class	Product Identification	Distribution	Affected Dates
I	Affected Lots: L40806 L40811 L40895 L40976 L41014 L41025 L41067 L41162 L41171 L41197 L41198 L41250	CA , nationwide	Manufactured and distributed from: December 2013 to March 2015

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/Safety/Recalls/ucm460169.htm>