



California Medical Device Recall Information



Recall Name

Insulet Corporation Recalls OmniPod Insulin Management System Due to Failure of the Needle Mechanism to Deploy

Recall Date	Product Description	Recalling Firm	Recall Reason
11/02/15	OmniPod Insulin Management System Catalog Number: <ul style="list-style-type: none"> • POD-ZXP420 	Insulet Corporation Billerica, MA	<i>Due to a slight increase in the reported cases in which the Pod's needle mechanism failed to deploy or there was a delay in the deployment of the needle mechanism.</i>
Recall Class	Product Identification	Distribution	Affected Dates
I	Affected Lots: L41880 L41901 L41881 L41902 L41892 L41903 L41895 L41904 L41897 L41905 L41898 L41906 L41899 L41907 L41900	CA , nationwide	Manufactured: July 2015 to August 2015 Distributed: September 2015

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm478915.htm>