



California Medical Device Recall Information



Recall Name

Conmed Corporation Recalls

PadPro and R2 Multi-function Defibrillation Electrodes

Due to Connector Compatibility with Specific Defibrillators

Recall Date	Product Description	Recalling Firm	Recall Reason
11/06/14	Conmed Multi-function Defibrillation Electrodes: <ul style="list-style-type: none"> • PadPro • R2 	Conmed Corporation Utica, NY	<i>These electrodes will not connect with Philips FR3 or FRx Defibrillator Units.</i>

Recall Class	Product Identification	Distribution	Affected Dates																		
I	All Lots <u>CONMED Catalog Numbers:</u> <table border="0" style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 20%;">2001H</td> <td>Adult Radiotransparent Electrode</td> </tr> <tr> <td>2001H-C</td> <td style="text-align: center;">“</td> </tr> <tr> <td>2001H-PC</td> <td style="text-align: center;">“</td> </tr> <tr> <td>2516H</td> <td style="text-align: center;">“</td> </tr> <tr> <td>2516H-PC</td> <td style="text-align: center;">“</td> </tr> <tr> <td>2603H</td> <td>Pediatric Radiotranslucent Electrode</td> </tr> <tr> <td>2602H</td> <td>Mini Pediatric Radiotranslucent Elec</td> </tr> <tr> <td>3115-1750</td> <td>Pediatric R2 Multifunction Electrode</td> </tr> <tr> <td>3115-1751</td> <td>R2 Multifunction Electrode</td> </tr> </table>	2001H	Adult Radiotransparent Electrode	2001H-C	“	2001H-PC	“	2516H	“	2516H-PC	“	2603H	Pediatric Radiotranslucent Electrode	2602H	Mini Pediatric Radiotranslucent Elec	3115-1750	Pediatric R2 Multifunction Electrode	3115-1751	R2 Multifunction Electrode	CA, nationwide	Distributed from: March 1, 2012 through October 29, 2014.
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FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm424871.htm>