



# California Medical Device Recall Information



## Recall Name

**GE Healthcare Recalls Single-Width Airway Modules (E-MiniC) and Accessories and Extension Modules (N-FC, N-FCREC) Due to a Potential for the CO2 Detector to Fail**

Recall Date	Product Description	Recalling Firm	Recall Reason
06/11/14	<ul style="list-style-type: none"> <li>Single-Width Airway Modules (E-miniC)</li> <li>Extension Modules N-FC and N-FCREC</li> <li>Modules serviced with FRU (Field Replaceable Unit)</li> </ul>	<b>GE Healthcare, LLC.</b> Waukesha, WI	<i>The affected CO2 detectors may fail or provide incorrect CO2 values for mechanically and spontaneous ventilated patients.</i>
Recall Class	Product Identification	Distribution	Affected Dates
I	<p><u>Single-Width Airway Modules (E-miniC)</u> Serial Numbers: <b>6818561 through 6898777</b></p> <p><u>Extension Modules N-FC and N-FCREC</u> Serial Numbers: <b>6799191 through 6905206</b></p> <p><u>FRU Catalog Number M1013204 (miniC Unit, NFCREC) Modules</u> serviced between: <b>February 2012 and May 2014</b></p>	<b>CA</b> , nationwide	<p>Manufacturing Dates: February 10, 2012 through October 2, 2012</p> <p>Distribution Dates: February 2012 to April 2014</p>

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm406734.htm>