



California Medical Device Recall Information



Recall Name

H&H Medical Corporation Recalls the Emergency Cricothyrotomy Kit Due to a Potential Defect

Recall Date	Product Description	Recalling Firm	Recall Reason																																												
8/27/13	H&H Emergency Cricothyrotomy Kit	H&H Medical Corp. Ordinary, VA	<i>Potential for a defective cuff balloon on the provided endotracheal airway.</i>																																												
Recall Class	Product Identification	Distribution	Affected Dates																																												
I	<table border="1"> <thead> <tr> <th>Lot #</th> <th>Expiration Date</th> </tr> </thead> <tbody> <tr><td>CKBD033</td><td>August 2015</td></tr> <tr><td>CKBE033</td><td>August 2015</td></tr> <tr><td>CKBD034</td><td>August 2015</td></tr> <tr><td>CKBF034</td><td>August 2015</td></tr> <tr><td>CKBG034</td><td>August 2015</td></tr> <tr><td>CKBP045</td><td>November 2015</td></tr> <tr><td>CKBP047</td><td>November 2015</td></tr> <tr><td>CKBQ047</td><td>November 2015</td></tr> <tr><td>CKBR060</td><td>February 2016</td></tr> <tr><td>CKBT065</td><td>April 2016</td></tr> <tr><td>CKBV070</td><td>May 2016</td></tr> <tr><td>CKBW070</td><td>May 2016</td></tr> <tr><td>CKBX070</td><td>May 2016</td></tr> <tr><td>CKBX071</td><td>May 2016</td></tr> <tr><td>CKBX076</td><td>June 2016</td></tr> <tr><td>CKBX078</td><td>July 2016</td></tr> <tr><td>CKBX079</td><td>July 2016</td></tr> <tr><td>CKBY079</td><td>July 2016</td></tr> <tr><td>CKBY080</td><td>July 2016</td></tr> <tr><td>CKBZ080</td><td>July 2016</td></tr> <tr><td>CKCA080</td><td>July 2016</td></tr> </tbody> </table>	Lot #	Expiration Date	CKBD033	August 2015	CKBE033	August 2015	CKBD034	August 2015	CKBF034	August 2015	CKBG034	August 2015	CKBP045	November 2015	CKBP047	November 2015	CKBQ047	November 2015	CKBR060	February 2016	CKBT065	April 2016	CKBV070	May 2016	CKBW070	May 2016	CKBX070	May 2016	CKBX071	May 2016	CKBX076	June 2016	CKBX078	July 2016	CKBX079	July 2016	CKBY079	July 2016	CKBY080	July 2016	CKBZ080	July 2016	CKCA080	July 2016	CA, nationwide	Produced between: August 16, 2012 and July 29, 2013
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FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/Safety/Recalls/ucm370775.htm>