



California Medical Device And Drug Recall Information



Recall Name

Nephron Pharmaceuticals Recalls EZ Breathe Atomizer Due to Possible Manufacturing Defect

Recall Date	Product Description	Recalling Firm	Recall Reason
4/30/13	<p>Sold as, or within...:</p> <ul style="list-style-type: none"> • EZ Breathe Atomizer Model #EZ-100 Device; or • Ashmanefrin Starter Kit (combo drug/device) NDC 0487-2784-10 	<p>Nephron Pharmaceuticals Corp. Orlando, FL</p>	<p><i>Potential for a quarter-inch diameter washer becoming dislodged, which could result in a choking hazard.</i></p>
Recall Class	Product Identification	Distribution	Affected Dates
I	<p>Affected Atomizer Serial Number Ranges (located on battery door):</p> <ul style="list-style-type: none"> • 1206034476 – 1206069065 • 1209069180 – 1209069202 • 1207003710 – 1207038299 • 1209069203 – 1209069460 • 1207046505 – 1207081124 • 1210000001 – 1210103680 • 1208027421 – 1208062155 • 1210104001 – 1210104044 • 1209000001 – 1209069179 <p>Suspect Kit Lots Recalled:</p> <ul style="list-style-type: none"> • R2029A • R2042A • R2029B • R2045A • R2039A • R2047A 	<p>CA, nationwide</p>	<p>Distributed from August 3, 2012 through April 24, 2013</p>

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm355016.htm>

<http://www.ezbreatheatomizer.com/2013/important-ez-breathe-atomizer-information.html>