

**K110136 RESPICHAMBER VALVED HOLDING CHAMBER**Apr 18, 2011  
90 days to decisionK110136 · Product code: **NVP** · Anesthesiology  
Source: <https://www.510kdatabase.net/k110136/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Holding Chambers, Direct Patient Interface (NVP)
Date received	Jan 18, 2011
Decision date	Apr 18, 2011
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Trudell Medical Intl.</b>
Location	London, CA
Contact	DARRYL FISHER
510(k) history	12 submissions · 12 cleared · 2002-2013

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k110136/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 24, 2026