

**K083883 CHARTIS CATHETER**Jun 25, 2009  
178 days to decisionK083883 · Product code: **CBI** · Anesthesiology  
Source: <https://www.510kdatabase.net/k083883/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tube, Tracheal/bronchial, Differential Ventilation (w/wo Connector) (CBI)
Date received	Dec 29, 2008
Decision date	Jun 25, 2009
Days to decision	178 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Pulmonx, Inc.</b>
Location	Palo Alto, CA, US
Contact	Roberta Hines
510(k) history	4 submissions · 4 cleared · 2009-2011

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k083883/> 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