

**K093888 COOPDECH ENDOBRONCHIAL BLOCKER TUBE**Mar 16, 2010  
88 days to decisionK093888 · Product code: **CBI** · Anesthesiology  
Source: <https://www.510kdatabase.net/k093888/>**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 18, 2009
Decision date	Mar 16, 2010
Days to decision	88 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Daiken Medical Co, Ltd.</b>
Location	Tokyo, JP
Contact	FUMIAKI KANAI
510(k) history	2 submissions · 2 cleared · 2008-2010

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k093888/> 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