

**K902687 STERI-CATH-DL**Aug 29, 1990  
71 days to decisionK902687 · Product code: **BSY** · Anesthesiology  
Source: <https://www.510kdatabase.net/k902687/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheters, Suction, Tracheobronchial (BSY)
Date received	Jun 19, 1990
Decision date	Aug 29, 1990
Days to decision	71 days
Third-party review	No

**APPLICANT**

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Company	<b>Concord/Portex</b>
Location	Keene, NH, US
Contact	ROBERT WHEELER
510(k) history	23 submissions · 20 cleared · 1989-1993

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k902687/>; 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 26, 2026