

K834565 ENDOBRONCHIAL TWIN LUMEN TUBEJan 17, 1984
20 days to decisionK834565 · Product code: **BTS** · AnesthesiologySource: <https://www.510kdatabase.net/k834565/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tube, Bronchial (w/wo Connector) (BTS)
Date received	Dec 28, 1983
Decision date	Jan 17, 1984
Days to decision	20 days
Third-party review	No

APPLICANT

Company	Portex, Inc.
Location	Walker, MI, US
510(k) history	20 submissions · 20 cleared · 1977-2004

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k834565/> Data sourced from FDA 510(k) public records (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. - Generated May 25, 2026