

K902116 MODIFIED ENDOBRONCHIAL TUBEAug 3, 1990
85 days to decisionK902116 · Product code: **BTS** · AnesthesiologySource: <https://www.510kdatabase.net/k902116/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tube, Bronchial (w/wo Connector) (BTS)
Date received	May 10, 1990
Decision date	Aug 3, 1990
Days to decision	85 days
Third-party review	No

APPLICANT

Company	Concord/Portex
Location	Keene, NH, US
Contact	EDWIN G GROVE
510(k) history	23 submissions · 20 cleared · 1989-1993

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k902116/>; 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 26, 2026