FDA 510(k) Premarket Notification

K902116 MODIFIED ENDOBRONCHIAL TUBE

CLEARED

Aug 3, 1990
85 days to decision

K902116 · Product code: **BTS** · Anesthesiology
Source: 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