FDA 510(k) Premarket Notification

K834565 ENDOBRONCHIAL TWIN LUMEN TUBE

CLEARED

Jan 17, 1984
20 days to decision

K834565 · Product code: **BTS** · Anesthesiology
Source: 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