FDA 510(k) Premarket Notification

K954707 BRONCHOSCOPE

CLEARED

Nov 22, 1995
41 days to decision

K954707 · Product code: **EOQ** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k954707/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bronchoscope (flexible Or Rigid) (EOQ)
Date received	Oct 12, 1995
Decision date	Nov 22, 1995
Days to decision	41 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Fujinon, Inc.**
Location	Washington, DC, US
Contact	GARY A ADLER
510(k) history	30 submissions · 30 cleared · 1990-2012