FDA 510(k) Premarket Notification

K832010 BRONCHO-FIBERSCOPE

CLEARED

Aug 1, 1983
39 days to decision

K832010 · Product code: **EOQ** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k832010/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bronchoscope (flexible Or Rigid) (EOQ)
Date received	Jun 23, 1983
Decision date	Aug 1, 1983
Days to decision	39 days
Third-party review	No

APPLICANT

Company	**KARL STORZ Endoscopy-America, Inc.**
Location	Mchenry, IL, US
510(k) history	361 submissions · 361 cleared · 1980-2024