FDA 510(k) Premarket Notification

K812039 SPIROMETER 210/211

CLEARED

Sep 25, 1981
66 days to decision

K812039 · Product code: **BZG** · Anesthesiology
Source: https://www.510kdatabase.net/k812039/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spirometer, Diagnostic (BZG)
Date received	Jul 21, 1981
Decision date	Sep 25, 1981
Days to decision	66 days
Third-party review	No

APPLICANT

Company	**Medix Medical Electronics (U.S.A.), Inc.**
Location	Mchenry, IL, US
510(k) history	14 submissions · 14 cleared · 1981-1986