FDA 510(k) Premarket Notification

K770141 INSPIROMETER RESPIRATORY EXERCISER

CLEARED

Feb 1, 1977
8 days to decision

K770141 · Product code: **BWF** · Anesthesiology
Source: https://www.510kdatabase.net/k770141/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spirometer, Therapeutic (incentive) (BWF)
Date received	Jan 24, 1977
Decision date	Feb 1, 1977
Days to decision	8 days
Third-party review	No

APPLICANT

Company	**Medi, Inc.**
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1977-1977