FDA 510(k) Premarket Notification

K832401 RESPIROMETER

CLEARED

Apr 30, 1984
285 days to decision

K832401 · Product code: **BZG** · Anesthesiology
Source: https://www.510kdatabase.net/k832401/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spirometer, Diagnostic (BZG)
Date received	Jul 20, 1983
Decision date	Apr 30, 1984
Days to decision	285 days
Third-party review	No

APPLICANT

Company	**Kinetix**
Location	Walker, MI, US
510(k) history	4 submissions · 4 cleared · 1983-1984