FDA 510(k) Premarket Notification

K833337 VOLUREX II

CLEARED

Nov 28, 1983
62 days to decision

K833337 · Product code: **BWF** · Anesthesiology
Source: https://www.510kdatabase.net/k833337/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spirometer, Therapeutic (incentive) (BWF)
Date received	Sep 27, 1983
Decision date	Nov 28, 1983
Days to decision	62 days
Third-party review	No

APPLICANT

Company	**Dhd Medical Products Div. Diemolding Corp.**
Location	Mchenry, IL, US
510(k) history	43 submissions · 43 cleared · 1979-1988