FDA 510(k) Premarket Notification

K982889 MEDPULSE RESPIRATORY VEST SYSTEM, MODEL 1000

CLEARED

Jun 1, 1999
288 days to decision

K982889 · Product code: **BYI** · Anesthesiology
Source: https://www.510kdatabase.net/k982889/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Percussor, Powered-electric (BYI)
Date received	Aug 17, 1998
Decision date	Jun 1, 1999
Days to decision	288 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Electromed, Inc.**
Location	New Prague, MN, US
Contact	CHET SIEVERT
510(k) history	5 submissions · 5 cleared · 1999-2022