

K982889 MEDPULSE RESPIRATORY VEST SYSTEM, MODEL 1000Jun 1, 1999
288 days to decisionK982889 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k982889/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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