FDA 510(k) Premarket Notification

K905489 VER-MED A10017 UNIVERSAL ELECTRODE

CLEARED

Mar 21, 1991
104 days to decision

K905489 · Product code: **DRX** · Cardiovascular
Source: https://www.510kdatabase.net/k905489/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Dec 7, 1990
Decision date	Mar 21, 1991
Days to decision	104 days
Third-party review	No

APPLICANT

Company	**Vermont Medical, Inc.**
Location	Bellows Falls, VT, US
Contact	DAVID LOVELL
510(k) history	9 submissions · 9 cleared · 1978-2003