FDA 510(k) Premarket Notification

K910139 VER-MED A10014 RESTING EKG ELECTRODE & A10021

CLEARED

Aug 15, 1991
213 days to decision

K910139 · Product code: **DRX** · Cardiovascular
Source: https://www.510kdatabase.net/k910139/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Jan 14, 1991
Decision date	Aug 15, 1991
Days to decision	213 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