FDA 510(k) Premarket Notification

K030073 VERMED A10021 RESTING EKG TAB ELECTRODE

CLEARED

Mar 28, 2003
79 days to decision

K030073 · Product code: **DRX** · Cardiovascular
Source: https://www.510kdatabase.net/k030073/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Jan 8, 2003
Decision date	Mar 28, 2003
Days to decision	79 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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