FDA 510(k) Premarket Notification

K090180 TENDER-TRODE PREWIRED PEDIATRIC ECG ELECTRODE

CLEARED

Jun 17, 2009
145 days to decision

K090180 · Product code: **DRX** · Cardiovascular
Source: https://www.510kdatabase.net/k090180/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Jan 23, 2009
Decision date	Jun 17, 2009
Days to decision	145 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Vermed, Inc.**
Location	Bellows Falls, VT, US
Contact	MARC FILLION
510(k) history	1 submissions · 1 cleared · 2009-2009