FDA 510(k) Premarket Notification

K971444 3M RED DOT 2234 & 2266 RADIOLUCENT MONITORING ELECTRODES

CLEARED

Jun 11, 1997
51 days to decision

K971444 · Product code: **DRX** · Cardiovascular
Source: https://www.510kdatabase.net/k971444/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Apr 21, 1997
Decision date	Jun 11, 1997
Days to decision	51 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**3M Medical Products Group**
Location	St Paul, MN, US
Contact	LINDA JOHNSEN
510(k) history	12 submissions · 12 cleared · 1991-1997