

**K971444 3M RED DOT 2234 & 2266 RADIOLUCENT  
MONITORING ELECTRODES**Jun 11, 1997  
51 days to decisionK971444 · Product code: **DRX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k971444/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k971444/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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