

**K933278 INTERMEDICS TRIANGLE CHEST PROBE MODEL
526-04**Mar 15, 1995
624 days to decisionK933278 · Product code: **KRG** · Cardiovascular
Source: <https://www.510kdatabase.net/k933278/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Programmer, Pacemaker (KRG)
Date received	Jun 29, 1993
Decision date	Mar 15, 1995
Days to decision	624 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Intermedics, Inc.
Location	Mchenry, IL, US
Contact	TODD A FONSECA
510(k) history	211 submissions · 201 cleared · 1977-1996

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k933278/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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