FDA 510(k) Premarket Notification

K933278 INTERMEDICS TRIANGLE CHEST PROBE MODEL 526-04

CLEARED

Mar 15, 1995
624 days to decision

K933278 · 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