FDA 510(k) Premarket Notification

K790220 PACEMAKER PROGRAMMER

CLEARED

Jul 31, 1979
182 days to decision

K790220 · Product code: **KRG** · Cardiovascular
Source: https://www.510kdatabase.net/k790220/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Programmer, Pacemaker (KRG)
Date received	Jan 30, 1979
Decision date	Jul 31, 1979
Days to decision	182 days
Third-party review	No

APPLICANT

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