FDA 510(k) Premarket Notification

K812813 INTERMEDICS PROGRAMMER, MODEL 522-03

CLEARED

Oct 27, 1981
21 days to decision

K812813 · Product code: **KRG** · Cardiovascular
Source: https://www.510kdatabase.net/k812813/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Programmer, Pacemaker (KRG)
Date received	Oct 6, 1981
Decision date	Oct 27, 1981
Days to decision	21 days
Third-party review	No

APPLICANT

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