FDA 510(k) Premarket Notification

K822246 NONINVASIVE TEMPORARY PACEMAKER

CLEARED

Aug 6, 1982
8 days to decision

K822246 · Product code: **DRO** · Cardiovascular
Source: https://www.510kdatabase.net/k822246/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pacemaker, Cardiac, External Transcutaneous (non-invasive) (DRO)
Date received	Jul 29, 1982
Decision date	Aug 6, 1982
Days to decision	8 days
Third-party review	No

APPLICANT

Company	**Ross Research, Inc.**
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1982-1984