FDA 510(k) Premarket Notification

K880738 MODEL 2331T PATIENT PROGRAMMER

CLEARED

Nov 17, 1988
267 days to decision

K880738 · Product code: **KRG** · Cardiovascular
Source: https://www.510kdatabase.net/k880738/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Programmer, Pacemaker (KRG)
Date received	Feb 24, 1988
Decision date	Nov 17, 1988
Days to decision	267 days
Third-party review	No

APPLICANT

Company	**Medtronic Vascular**
Location	Walker, MI, US
Contact	H SWANSON
510(k) history	475 submissions · 453 cleared · 1977-2023