FDA 510(k) Premarket Notification

K781327 PULSE GENERATOR, MODEL 240, EXTERNAL

CLEARED

Dec 11, 1978
132 days to decision

K781327 · Product code: **DTE** · Cardiovascular
Source: https://www.510kdatabase.net/k781327/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pulse-generator, Pacemaker, External (DTE)
Date received	Aug 1, 1978
Decision date	Dec 11, 1978
Days to decision	132 days
Third-party review	No

APPLICANT

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