FDA 510(k) Premarket Notification

K802040 BATTERY FOR #229-01 PULSE GENERATOR

CLEARED

Sep 9, 1980
29 days to decision

K802040 · Product code: **DXY** · Cardiovascular
Source: https://www.510kdatabase.net/k802040/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Aug 11, 1980
Decision date	Sep 9, 1980
Days to decision	29 days
Third-party review	No

APPLICANT

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