

K781327 PULSE GENERATOR, MODEL 240, EXTERNALDec 11, 1978
132 days to decisionK781327 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k781327/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 28, 2026