

K791093 MODELS 2411/2418 VENTRICULAR-INHIBITEDJun 20, 1979
8 days to decisionK791093 · Product code: **DXY** · CardiovascularSource: <https://www.510kdatabase.net/k791093/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Jun 12, 1979
Decision date	Jun 20, 1979
Days to decision	8 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k791093/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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