

K770211 MICROPULSE MODEL 20SMar 7, 1977
35 days to decisionK770211 · Product code: **DXY** · CardiovascularSource: <https://www.510kdatabase.net/k770211/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Jan 31, 1977
Decision date	Mar 7, 1977
Days to decision	35 days
Third-party review	No

APPLICANT

Company	Edwards Pacemaker Systems
Location	Mchenry, IL, US
510(k) history	13 submissions · 13 cleared · 1977-1979

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Device record: <https://www.510kdatabase.net/k770211/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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