

K782146 PACEMAKER, MODEL DDP ELECTRONIC CIRCUIT

Dec 27, 1978

K782146 · Product code: **DXY** · CardiovascularSource: <https://www.510kdatabase.net/k782146/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent - SP
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Dec 27, 1978
Decision date	Dec 27, 1978
Third-party review	No

APPLICANT

Company	Arco Medical Products Co.
Location	Walker, MI, US
510(k) history	21 submissions · 20 cleared · 1976-1980

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Device record: <https://www.510kdatabase.net/k782146/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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