FDA 510(k) Premarket Notification

K782146 PACEMAKER, MODEL DDP ELECTRONIC CIRCUIT

CLEARED

Dec 27, 1978

K782146 · Product code: **DXY** · Cardiovascular
Source: 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