

K791715 ARCOLITH 2000Sep 27, 1979
27 days to decisionK791715 · Product code: **DXY** · CardiovascularSource: <https://www.510kdatabase.net/k791715/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Aug 31, 1979
Decision date	Sep 27, 1979
Days to decision	27 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k791715/> 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 July 9, 2026