

K780430 LITHICRON F PROGRAMMABLE PULSE GENDec 22, 1978
280 days to decisionK780430 · Product code: **DXY** · CardiovascularSource: <https://www.510kdatabase.net/k780430/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Mar 17, 1978
Decision date	Dec 22, 1978
Days to decision	280 days
Third-party review	No

APPLICANT

Company	Medcor, Inc.
Location	Mchenry, IL, US
510(k) history	8 submissions · 6 cleared · 1977-1980

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k780430/> 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