FDA 510(k) Premarket Notification

K780430 LITHICRON F PROGRAMMABLE PULSE GEN

CLEARED

Dec 22, 1978
280 days to decision

K780430 · Product code: **DXY** · Cardiovascular
Source: 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