FDA 510(k) Premarket Notification

K770559 C-L0L CARDIAC PACEMAKER

CLEARED

Mar 28, 1977
5 days to decision

K770559 · Product code: **DXY** · Cardiovascular
Source: https://www.510kdatabase.net/k770559/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Mar 23, 1977
Decision date	Mar 28, 1977
Days to decision	5 days
Third-party review	No

APPLICANT

Company	**Coratomic, Inc.**
Location	Mchenry, IL, US
510(k) history	15 submissions · 14 cleared · 1976-1986