FDA 510(k) Premarket Notification

K771992 L-501 CARDIAC PACEMAKER

CLEARED

Nov 2, 1977
12 days to decision

K771992 · Product code: **DXY** · Cardiovascular
Source: https://www.510kdatabase.net/k771992/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Oct 21, 1977
Decision date	Nov 2, 1977
Days to decision	12 days
Third-party review	No

APPLICANT

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