FDA 510(k) Premarket Notification

K791419 OMNICOR PROGRAMMER MODEL 222C

CLEARED

Nov 27, 1979
133 days to decision

K791419 · Product code: **KRG** · Cardiovascular
Source: https://www.510kdatabase.net/k791419/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Programmer, Pacemaker (KRG)
Date received	Jul 17, 1979
Decision date	Nov 27, 1979
Days to decision	133 days
Third-party review	No

APPLICANT

Company	**Cordis Corp.**
Location	Mchenry, IL, US
Website	https://cordis.com
510(k) history	315 submissions · 281 cleared · 1976-2014

Cordis Corp. is a medical device manufacturer based in McHenry, US. The company specializes in interventional cardiovascular and gastroenterology devices. Cordis has a substantial FDA 510(k) regulatory history spanning from 1976 to 2014. The company received FDA 510(k) clearances from total submissions. Its portfolio focuses primarily on cardiovascular devices and gastroenterology stent systems, including percutaneous transluminal angioplasty catheters, emboli capture guidewires, and self-expanding biliary stent systems. Notable cleared products include the FLEXSTENT Bili...

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k791419/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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