FDA 510(k) Premarket Notification

K841232 PULSE GENERATOR 696, 696B & 696L

CLEARED

Dec 3, 1984
256 days to decision

K841232 · Product code: **DXY** · Cardiovascular
Source: https://www.510kdatabase.net/k841232/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Mar 22, 1984
Decision date	Dec 3, 1984
Days to decision	256 days
Third-party review	No

APPLICANT

Company	**Siemens Elema AB**
Location	Mchenry, IL, US
510(k) history	63 submissions · 60 cleared · 1978-2003