FDA 510(k) Premarket Notification

K792223 PACESETTER MODEL 350

CLEARED

Nov 30, 1979
25 days to decision

K792223 · Product code: **DTE** · Cardiovascular
Source: https://www.510kdatabase.net/k792223/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pulse-generator, Pacemaker, External (DTE)
Date received	Nov 5, 1979
Decision date	Nov 30, 1979
Days to decision	25 days
Third-party review	No

APPLICANT

Company	**Cynergy**
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1978-1981