FDA 510(k) Premarket Notification

K770865 MONITOR, PACEMAKER

CLEARED

Jun 14, 1977
34 days to decision

K770865 · Product code: **DTC** · Cardiovascular
Source: https://www.510kdatabase.net/k770865/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Pacemaker Generator Function (DTC)
Date received	May 11, 1977
Decision date	Jun 14, 1977
Days to decision	34 days
Third-party review	No

APPLICANT

Company	**Adcor Electronics, Inc.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1977-1977