FDA 510(k) Premarket Notification

K801799 ATRIAL/VENTRICULAR SEQENTIAL #080-20000

CLEARED

Jan 2, 1981
157 days to decision

K801799 · Product code: **LDF** · Cardiovascular
Source: https://www.510kdatabase.net/k801799/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Jul 29, 1980
Decision date	Jan 2, 1981
Days to decision	157 days
Third-party review	No

APPLICANT

Company	**Medical Testing System, Inc.**
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1977-1981