FDA 510(k) Premarket Notification

K802259 TRANSVENOUS PACEMAKER ELECTRODE KIT

CLEARED

Oct 23, 1980
37 days to decision

K802259 · Product code: **LDF** · Cardiovascular
Source: https://www.510kdatabase.net/k802259/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Sep 16, 1980
Decision date	Oct 23, 1980
Days to decision	37 days
Third-party review	No

APPLICANT

Company	**Stanco Medical, Inc.**
Location	Mchenry, IL, US
510(k) history	19 submissions · 19 cleared · 1979-1981

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k802259/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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