FDA 510(k) Premarket Notification

CLEARED

K792604 LIFELINE TLE ENDOCARDIAL PACING LEAD,

Dec 31, 1979
13 days to decision

K792604 · Product code: **DTB** · Cardiovascular
Source: https://www.510kdatabase.net/k792604/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Dec 18, 1979
Decision date	Dec 31, 1979
Days to decision	13 days
Third-party review	No

APPLICANT

Company	**Intermedics, Inc.**
Location	Mchenry, IL, US
510(k) history	211 submissions · 201 cleared · 1977-1996