FDA 510(k) Premarket Notification

K832110 ELECTRODE LEADS 030-267 & 030-268

CLEARED

Aug 12, 1983
43 days to decision

K832110 · Product code: **DTB** · Cardiovascular
Source: https://www.510kdatabase.net/k832110/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Jun 30, 1983
Decision date	Aug 12, 1983
Days to decision	43 days
Third-party review	No

APPLICANT

Company	**Telectronics, Inc.**
Location	Mchenry, IL, US
510(k) history	107 submissions · 107 cleared · 1977-1990