

K832110 ELECTRODE LEADS 030-267 & 030-268Aug 12, 1983
43 days to decisionK832110 · Product code: **DTB** · CardiovascularSource: <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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k832110/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 22, 2026