

K800354 LEAD, TINED PACING, IMPLANTABLEMay 30, 1980
105 days to decisionK800354 · Product code: **DTB** · CardiovascularSource: <https://www.510kdatabase.net/k800354/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Feb 15, 1980
Decision date	May 30, 1980
Days to decision	105 days
Third-party review	No

APPLICANT

Company	Daig Corp.
Location	Mchenry, IL, US
510(k) history	63 submissions · 63 cleared · 1977-2000

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k800354/> 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 July 3, 2026