

K812270 PERM. ENDOCARDIAL UNIPOLAR TINED ATRIALSep 16, 1981
34 days to decisionK812270 · Product code: **DTB** · CardiovascularSource: <https://www.510kdatabase.net/k812270/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Aug 13, 1981
Decision date	Sep 16, 1981
Days to decision	34 days
Third-party review	No

APPLICANT

Company	Diag Corp.
Location	Walker, MI, US
510(k) history	9 submissions · 9 cleared · 1980-1999

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k812270/> 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 4, 2026