

K820121 LASERPOR AP ATRIAL ENDOCARD. ELECTR. LD.Feb 12, 1982
25 days to decisionK820121 · Product code: **DTB** · CardiovascularSource: <https://www.510kdatabase.net/k820121/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Jan 18, 1982
Decision date	Feb 12, 1982
Days to decision	25 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.netDevice record: <https://www.510kdatabase.net/k820121/> 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