FDA 510(k) Premarket Notification

K820121 LASERPOR AP ATRIAL ENDOCARD. ELECTR. LD.

CLEARED

Feb 12, 1982
25 days to decision

K820121 · Product code: **DTB** · Cardiovascular
Source: 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