FDA 510(k) Premarket Notification

K802552 PERM. ENDOCARDIAL UNIPOLAR TINED/WEDGE

CLEARED

Jan 16, 1981
93 days to decision

K802552 · Product code: **DTB** · Cardiovascular
Source: https://www.510kdatabase.net/k802552/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Oct 15, 1980
Decision date	Jan 16, 1981
Days to decision	93 days
Third-party review	No

APPLICANT

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