FDA 510(k) Premarket Notification

K781620 PACING LEAD MODELS, MDM 1060, MDM 1090

CLEARED

Dec 8, 1978
79 days to decision

K781620 · Product code: **DTB** · Cardiovascular
Source: https://www.510kdatabase.net/k781620/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Sep 20, 1978
Decision date	Dec 8, 1978
Days to decision	79 days
Third-party review	No

APPLICANT

Company	**R.E. Brown Co., Inc.**
Location	Mchenry, IL, US
510(k) history	10 submissions · 10 cleared · 1978-1978