

K781620 PACING LEAD MODELS, MDM 1060, MDM 1090Dec 8, 1978
79 days to decisionK781620 · Product code: **DTB** · CardiovascularSource: <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

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Device record: <https://www.510kdatabase.net/k781620/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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