

**K781623 PACING MODELS MUM 860, MUM 890**Dec 8, 1978  
79 days to decisionK781623 · Product code: **DTB** · Cardiovascular  
Source: <https://www.510kdatabase.net/k781623/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k781623/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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