

**K800891 RESTERILIZATION OF IMPLANTABLE ELECTRODE**May 28, 1980  
40 days to decisionK800891 · Product code: **DTB** · CardiovascularSource: <https://www.510kdatabase.net/k800891/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Apr 18, 1980
Decision date	May 28, 1980
Days to decision	40 days
Third-party review	No

**APPLICANT**

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Company	<b>Telectronics, Inc.</b>
Location	Mchenry, IL, US
510(k) history	107 submissions · 107 cleared · 1977-1990

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

Device record: <https://www.510kdatabase.net/k800891/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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