

**K870610 RESUBMISSION OF MODEL 030-437 IMPLANT.  
ELECTRODE**Mar 20, 1987  
46 days to decisionK870610 · Product code: **DTB** · Cardiovascular  
Source: <https://www.510kdatabase.net/k870610/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Feb 2, 1987
Decision date	Mar 20, 1987
Days to decision	46 days
Third-party review	No

**APPLICANT**

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Company	<b>Telectronics, Inc.</b>
Location	Mchenry, IL, US
Contact	WILLIAM C NEALON
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/k870610/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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