

**K031210 MODEL 4951M MYOCARDIAL UNIPOLAR LEAD**May 16, 2003  
29 days to decisionK031210 · Product code: **DTB** · CardiovascularSource: <https://www.510kdatabase.net/k031210/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Apr 17, 2003
Decision date	May 16, 2003
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Metronic, Inc.</b>
Location	Mchenry, IL, US
Contact	TINA BENOIT
510(k) history	4 submissions · 4 cleared · 1983-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k031210/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 15, 2026