

**K810928 ENDOCARDIAL ELECTRODE**Jul 2, 1981  
87 days to decisionK810928 · Product code: **DTB** · CardiovascularSource: <https://www.510kdatabase.net/k810928/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Apr 6, 1981
Decision date	Jul 2, 1981
Days to decision	87 days
Third-party review	No

**APPLICANT**

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Company	<b>Vitatron Medical BV</b>
Location	Mchenry, IL, US
510(k) history	19 submissions · 19 cleared · 1976-1986

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

Device record: <https://www.510kdatabase.net/k810928/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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