

K791732 IMPLANTABLE ENDOCARDIAL SCREW-IN ELECSep 19, 1979
14 days to decisionK791732 · Product code: **DTB** · CardiovascularSource: <https://www.510kdatabase.net/k791732/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Sep 5, 1979
Decision date	Sep 19, 1979
Days to decision	14 days
Third-party review	No

APPLICANT

Company	Oscor, Inc.
Location	Palm Harbor, FL, US
510(k) history	49 submissions · 46 cleared · 1979-2021

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Device record: <https://www.510kdatabase.net/k791732/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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