

K831926 IMPLANTABLE ENDOCARDIAL PACING LEADSMar 12, 1984
271 days to decisionK831926 · Product code: **DTG** · CardiovascularSource: <https://www.510kdatabase.net/k831926/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Magnet, Test, Pacemaker (DTG)
Date received	Jun 15, 1983
Decision date	Mar 12, 1984
Days to decision	271 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/k831926/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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