FDA 510(k) Premarket Notification

K791732 IMPLANTABLE ENDOCARDIAL SCREW-IN ELEC

CLEARED

Sep 19, 1979
14 days to decision

K791732 · Product code: **DTB** · Cardiovascular
Source: 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