FDA 510(k) Premarket Notification

K830950 VASCULAR PRODUCTS IMPLANTABLE PACING

CLEARED

Apr 18, 1983
24 days to decision

K830950 · Product code: **DTB** · Cardiovascular
Source: https://www.510kdatabase.net/k830950/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Mar 25, 1983
Decision date	Apr 18, 1983
Days to decision	24 days
Third-party review	No

APPLICANT

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