FDA 510(k) Premarket Notification

K920662 VASCOTWIST 9/60

CLEARED

Nov 24, 1993
649 days to decision

K920662 · Product code: **DTB** · Cardiovascular
Source: https://www.510kdatabase.net/k920662/

SUBMISSION DETAILS

Decision	Substantially Equivalent - SP
Submission type	Traditional
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Feb 14, 1992
Decision date	Nov 24, 1993
Days to decision	649 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Vascor Medical Corp.**
Location	Tarpon Springs, FL, US
Contact	AUDREY MACCIA
510(k) history	3 submissions · 2 cleared · 1993-1993

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k920662/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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