FDA 510(k) Premarket Notification

K901780 RTM 36-05, 60-05, RTD 36-05, 60-05 RADIO. LEADWIRE

CLEARED

Oct 9, 1990
175 days to decision

K901780 · Product code: **DSA** · Cardiovascular
Source: https://www.510kdatabase.net/k901780/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cable, Transducer And Electrode, Patient, (including Connector) (DSA)
Date received	Apr 17, 1990
Decision date	Oct 9, 1990
Days to decision	175 days
Third-party review	No

APPLICANT

Company	**Medtronic Vascular**
Location	Walker, MI, US
Contact	JANICE M PEVIDE
510(k) history	475 submissions · 453 cleared · 1977-2023