FDA 510(k) Premarket Notification

K900519 CLASSIC ECG/STIMULATING NO. 1700 SERIES

CLEARED

Mar 19, 1990
45 days to decision

K900519 · Product code: **DRX** · Cardiovascular
Source: https://www.510kdatabase.net/k900519/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Feb 2, 1990
Decision date	Mar 19, 1990
Days to decision	45 days
Third-party review	No

APPLICANT

Company	**Classic Medical Products, Inc.**
Location	Muskego, WI, US
Contact	ROBERT A MACUR
510(k) history	2 submissions · 2 cleared · 1986-1990