FDA 510(k) Premarket Notification

K896569 ELECTROTRACE DISPOSABLE AG/AGCL EKG MONITOR ELEC.

CLEARED

Jan 18, 1990
59 days to decision

K896569 · Product code: **DRX** · Cardiovascular
Source: https://www.510kdatabase.net/k896569/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Nov 20, 1989
Decision date	Jan 18, 1990
Days to decision	59 days
Third-party review	No

APPLICANT

Company	**Jason Marketing Co.**
Location	Huntington Beach, CA, US
Contact	JAY RUEHLEN
510(k) history	5 submissions · 5 cleared · 1987-1992