FDA 510(k) Premarket Notification

K933603 LEADWIRE

CLEARED

Jan 18, 1994
175 days to decision

K933603 · Product code: **DSA** · Cardiovascular
Source: https://www.510kdatabase.net/k933603/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cable, Transducer And Electrode, Patient, (including Connector) (DSA)
Date received	Jul 27, 1993
Decision date	Jan 18, 1994
Days to decision	175 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Sentry Medical Products, Inc.**
Location	Walker, MI, US
Contact	WILLIAM E BLAIR
510(k) history	13 submissions · 13 cleared · 1983-1996