FDA 510(k) Premarket Notification

K932704 PATIENT CABLE

CLEARED

Dec 27, 1993
207 days to decision

K932704 · Product code: **DSA** · Cardiovascular
Source: https://www.510kdatabase.net/k932704/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cable, Transducer And Electrode, Patient, (including Connector) (DSA)
Date received	Jun 3, 1993
Decision date	Dec 27, 1993
Days to decision	207 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