

K932704 PATIENT CABLEDec 27, 1993
207 days to decisionK932704 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k932704/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 1, 2026