

K896699 TYPE 9040/NEOApr 20, 1990
143 days to decisionK896699 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k896699/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Nov 28, 1989
Decision date	Apr 20, 1990
Days to decision	143 days
Third-party review	No

APPLICANT

Company	S & W Medico Teknik
Location	Walker, MI, US
Contact	LISBETH ISBRANDT
510(k) history	46 submissions · 46 cleared · 1984-1995

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k896699/>; 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 May 26, 2026