

K896386 MINIMON 7133B PATIENT MONITORDec 13, 1989
37 days to decisionK896386 · Product code: **DPS** · CardiovascularSource: <https://www.510kdatabase.net/k896386/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	Nov 6, 1989
Decision date	Dec 13, 1989
Days to decision	37 days
Third-party review	No

APPLICANT

Company	Kontron Instruments, Inc.
Location	Mchenry, IL, US
Contact	DAVID CROMWICK
510(k) history	57 submissions · 57 cleared · 1981-1993

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k896386/>; 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 21, 2026