

K890791 SIEMENS SIRECUST 730 PATIENT MONITORMay 2, 1989
74 days to decisionK890791 · Product code: **DPS** · CardiovascularSource: <https://www.510kdatabase.net/k890791/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	Feb 17, 1989
Decision date	May 2, 1989
Days to decision	74 days
Third-party review	No

APPLICANT

Company	Siemens Medical Electronics
Location	Danvers, MA, US
Contact	MURFITT, PHD
510(k) history	15 submissions · 15 cleared · 1988-1995

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k890791/>, 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