

K881770 EPICARDIA LE/FDMay 31, 1988
35 days to decisionK881770 · Product code: **DQK** · CardiovascularSource: <https://www.510kdatabase.net/k881770/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Computer, Diagnostic, Programmable (DQK)
Date received	Apr 26, 1988
Decision date	May 31, 1988
Days to decision	35 days
Third-party review	No

APPLICANT

Company	Medicomp, Inc.
Location	Mchenry, IL, US
Contact	JOE RIFFE
510(k) history	23 submissions · 23 cleared · 1983-2018

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k881770/>, 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