FDA 510(k) Premarket Notification

K881770 EPICARDIA LE/FD

CLEARED

May 31, 1988
35 days to decision

K881770 · Product code: **DQK** · Cardiovascular
Source: 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