

K800092 COMBISON COMPUTERApr 8, 1980
85 days to decisionK800092 · Product code: **DQK** · CardiovascularSource: <https://www.510kdatabase.net/k800092/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Computer, Diagnostic, Programmable (DQK)
Date received	Jan 14, 1980
Decision date	Apr 8, 1980
Days to decision	85 days
Third-party review	No

APPLICANT

Company	Kt Medical, Inc.
Location	Mchenry, IL, US
510(k) history	13 submissions · 13 cleared · 1980-1980

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Device record: <https://www.510kdatabase.net/k800092/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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