

**K810731 MODEL IM1000 INTEGRATED MONITOR**Mar 27, 1981  
10 days to decisionK810731 · Product code: **DQK** · CardiovascularSource: <https://www.510kdatabase.net/k810731/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Computer, Diagnostic, Programmable (DQK)
Date received	Mar 17, 1981
Decision date	Mar 27, 1981
Days to decision	10 days
Third-party review	No

**APPLICANT**

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Company	<b>Gould, Inc.</b>
Location	Mchenry, IL, US
510(k) history	31 submissions · 31 cleared · 1976-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k810731/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated July 9, 2026