FDA 510(k) Premarket Notification

K810731 MODEL IM1000 INTEGRATED MONITOR

CLEARED

Mar 27, 1981
10 days to decision

K810731 · Product code: **DQK** · Cardiovascular
Source: https://www.510kdatabase.net/k810731/

SUBMISSION DETAILS

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

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