

K830885 GOULD MODEL IM3000-IM3100 BATTERY POWERJun 22, 1983
93 days to decisionK830885 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k830885/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Mar 21, 1983
Decision date	Jun 22, 1983
Days to decision	93 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k830885/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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