

K820540 OLLI 565Apr 1, 1982
30 days to decisionK820540 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k820540/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Mar 2, 1982
Decision date	Apr 1, 1982
Days to decision	30 days
Third-party review	No

APPLICANT

Company	Kone Instruments, Inc.
Location	Mchenry, IL, US
510(k) history	12 submissions · 12 cleared · 1982-1987

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Device record: <https://www.510kdatabase.net/k820540/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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