

K790644 SUBMAXApr 26, 1979
27 days to decisionK790644 · Product code: **DRT** · Cardiovascular
Source: <https://www.510kdatabase.net/k790644/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Mar 30, 1979
Decision date	Apr 26, 1979
Days to decision	27 days
Third-party review	No

APPLICANT

Company	Chase Mfg. Co.
Location	Mchenry, IL, US
510(k) history	10 submissions · 10 cleared · 1977-1979

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Device record: <https://www.510kdatabase.net/k790644/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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