

K844348 DYNASCOPE MONITOR DS-800Dec 4, 1984
35 days to decisionK844348 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k844348/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Oct 30, 1984
Decision date	Dec 4, 1984
Days to decision	35 days
Third-party review	No

APPLICANT

Company	Fukuda Denshi USA, Inc.
Location	Mchenry, IL, US
Contact	ROBERT J STEURER
510(k) history	68 submissions · 68 cleared · 1984-2018

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k844348/>; Data sourced from FDA 510(k) public records (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. - Generated May 18, 2026