

K935817 MEDSIM 300 PATIENT SIMULATORJan 18, 1994
47 days to decisionK935817 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k935817/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Dec 2, 1993
Decision date	Jan 18, 1994
Days to decision	47 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Dynatech/Nevada, Inc.
Location	Carson City, NV, US
Contact	HOWARD M HOLSTEIN
510(k) history	6 submissions · 6 cleared · 1990-1995

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