FDA 510(k) Premarket Notification

K897074 MODEL 215M PATIENT SIMULATOR

CLEARED

May 17, 1990
146 days to decision

K897074 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k897074/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Dec 22, 1989
Decision date	May 17, 1990
Days to decision	146 days
Third-party review	No

APPLICANT

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