

**K897074 MODEL 215M PATIENT SIMULATOR**May 17, 1990  
146 days to decisionK897074 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k897074/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k897074/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 28, 2026