

K891693 AUTOMATED THERAPY SYSTEMAug 21, 1989
152 days to decisionK891693 · Product code: **HCC** · Neurology
Source: <https://www.510kdatabase.net/k891693/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biofeedback (HCC)
Date received	Mar 22, 1989
Decision date	Aug 21, 1989
Days to decision	152 days
Third-party review	No

APPLICANT

Company	Harding & Harris, Inc.
Location	Orem, UT, US
Contact	DOUGLAS HARDING
510(k) history	1 submissions · 1 cleared · 1989-1989

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k891693/>; 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 July 1, 2026