FDA 510(k) Premarket Notification

K891693 AUTOMATED THERAPY SYSTEM

CLEARED

Aug 21, 1989
152 days to decision

K891693 · 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