FDA 510(k) Premarket Notification

K875308 PROFESSIONAL SERIES BIOFEEDBACK DEVICES MODEL 421

CLEARED

Feb 26, 1988
59 days to decision

K875308 · Product code: **HCC** · Neurology
Source: https://www.510kdatabase.net/k875308/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biofeedback (HCC)
Date received	Dec 29, 1987
Decision date	Feb 26, 1988
Days to decision	59 days
Third-party review	No

APPLICANT

Company	**Self Regulation Systems, Inc.**
Location	Walker, MI, US
Contact	CARL R CHRISTINE
510(k) history	3 submissions · 3 cleared · 1984-1988