FDA 510(k) Premarket Notification

K884852 BIOFEEDBACK DEVICE EMG1, EMG ULTRA LOW NOISE OPTI.

CLEARED

Feb 2, 1989
76 days to decision

K884852 · Product code: **HCC** · Neurology
Source: https://www.510kdatabase.net/k884852/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biofeedback (HCC)
Date received	Nov 18, 1988
Decision date	Feb 2, 1989
Days to decision	76 days
Third-party review	No

APPLICANT

Company	**Moe**
Location	Soquel, CA, US
Contact	MIKE WILBER
510(k) history	6 submissions · 6 cleared · 1989-1989