FDA 510(k) Premarket Notification

K832122 ENRAF NONIVS2 CHANNEL EMG/TEMP. FEED

CLEARED

Aug 12, 1983
43 days to decision

K832122 · Product code: **HCC** · Neurology
Source: https://www.510kdatabase.net/k832122/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biofeedback (HCC)
Date received	Jun 30, 1983
Decision date	Aug 12, 1983
Days to decision	43 days
Third-party review	No

APPLICANT

Company	**Fluidotherapy Corp.**
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1983-1985