

K832122 ENRAF NONIVS2 CHANNEL EMG/TEMP. FEEDAug 12, 1983
43 days to decisionK832122 · 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

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Device record: <https://www.510kdatabase.net/k832122/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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