

K801670 NDI-IKE EMGAug 4, 1980
13 days to decisionK801670 · Product code: **GWK** · Neurology
Source: <https://www.510kdatabase.net/k801670/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Conditioner, Signal, Physiological (GWK)
Date received	Jul 22, 1980
Decision date	Aug 4, 1980
Days to decision	13 days
Third-party review	No

APPLICANT

Company	Neurodiagnostics, Inc.
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1980-1984

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Device record: <https://www.510kdatabase.net/k801670/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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