

K802887 MODEL EEG 4221Dec 16, 1980
29 days to decisionK802887 · Product code: **GWQ** · Neurology
Source: <https://www.510kdatabase.net/k802887/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Full-montage Standard Electroencephalograph (GWQ)
Date received	Nov 17, 1980
Decision date	Dec 16, 1980
Days to decision	29 days
Third-party review	No

APPLICANT

Company	Nihon Kohden America, Inc.
Location	Foothill Ranch, CA, US
510(k) history	166 submissions · 163 cleared · 1979-2012

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Device record: <https://www.510kdatabase.net/k802887/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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