

K801604 PATHFINDER ELECTRODIAGNOSTIC SYSTEMAug 4, 1980
21 days to decisionK801604 · Product code: **GWK** · Neurology
Source: <https://www.510kdatabase.net/k801604/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Conditioner, Signal, Physiological (GWK)
Date received	Jul 14, 1980
Decision date	Aug 4, 1980
Days to decision	21 days
Third-party review	No

APPLICANT

Company	Nicolet Biomedical Instruments
Location	Mchenry, IL, US
510(k) history	26 submissions · 26 cleared · 1977-1988

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Device record: <https://www.510kdatabase.net/k801604/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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