

K800648 CADWELL 7200Apr 21, 1980
28 days to decisionK800648 · Product code: **IKN** · Neurology
Source: <https://www.510kdatabase.net/k800648/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electromyograph, Diagnostic (IKN)
Date received	Mar 24, 1980
Decision date	Apr 21, 1980
Days to decision	28 days
Third-party review	No

APPLICANT

Company	Cadwell Laboratories, Inc.
Location	Walker, MI, US
510(k) history	46 submissions · 46 cleared · 1979-2007

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Device record: <https://www.510kdatabase.net/k800648/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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