

K802637 TECA MODEL-10 ELECTROMYOGRAPHNov 26, 1980
35 days to decisionK802637 · Product code: **IKN** · Neurology
Source: <https://www.510kdatabase.net/k802637/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electromyograph, Diagnostic (IKN)
Date received	Oct 22, 1980
Decision date	Nov 26, 1980
Days to decision	35 days
Third-party review	No

APPLICANT

Company	Teca, Inc.
Location	Mchenry, IL, US
510(k) history	29 submissions · 29 cleared · 1976-1996

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Device record: <https://www.510kdatabase.net/k802637/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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