

K840321 ENDODYNE DUALMar 16, 1984
51 days to decisionK840321 · Product code: **LIH** · Neurology
Source: <https://www.510kdatabase.net/k840321/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Interferential Current Therapy (LIH)
Date received	Jan 25, 1984
Decision date	Mar 16, 1984
Days to decision	51 days
Third-party review	No

APPLICANT

Company	Elmed, Inc.
Location	Mchenry, IL, US
510(k) history	26 submissions · 26 cleared · 1977-2001

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Device record: <https://www.510kdatabase.net/k840321/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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