FDA 510(k) Premarket Notification

K840321 ENDODYNE DUAL

CLEARED

Mar 16, 1984
51 days to decision

K840321 · 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