FDA 510(k) Premarket Notification

K822396 THE 3500

CLEARED

Sep 24, 1982
45 days to decision

K822396 · Product code: **GWF** · Neurology
Source: https://www.510kdatabase.net/k822396/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Electrical, Evoked Response (GWF)
Date received	Aug 10, 1982
Decision date	Sep 24, 1982
Days to decision	45 days
Third-party review	No

APPLICANT

Company	**Tracor Northern, Inc.**
Location	Mchenry, IL, US
510(k) history	10 submissions · 10 cleared · 1981-1988

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k822396/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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