FDA 510(k) Premarket Notification

K802390 EMG-UT 30

CLEARED

Jan 9, 1981
99 days to decision

K802390 · Product code: **HCC** · Neurology
Source: https://www.510kdatabase.net/k802390/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biofeedback (HCC)
Date received	Oct 2, 1980
Decision date	Jan 9, 1981
Days to decision	99 days
Third-party review	No

APPLICANT

Company	**American Scientific Corp.**
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1981-1981