FDA 510(k) Premarket Notification

K094054 MODIFCATION TO: NUVASIVE NEUROVISION EMG ENDOTRACHEAL TUBE

CLEARED

May 14, 2010
134 days to decision

K094054 · Product code: **GWF** · Neurology
Source: https://www.510kdatabase.net/k094054/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Stimulator, Electrical, Evoked Response (GWF)
Date received	Dec 31, 2009
Decision date	May 14, 2010
Days to decision	134 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Nuvasive, Inc.**
Location	San Diego, CA, US
Contact	Sheila Bruschi
Website	http://www.nuvasive.com/
510(k) history	91 submissions · 90 cleared · 1999-2024

NuVasive, Inc. is a medical device company headquartered in San Diego, California. The company develops and markets surgical solutions focused on spine and orthopedic procedures. NuVasive operates globally and serves healthcare professionals and patients worldwide. The company maintains a strong FDA 510(k) regulatory record with FDA 510(k) clearances from total submissions since 1999. Orthopedic devices represent the dominant category, accounting for the majority of the company's cleared submissions. The most recent clearance was granted in 2024, demonstrating continued r...