

K013215 MODIFIED INS-1 INTRAOPERATIVE NERVE SURVEILLANCE SYSTEMOct 16, 2001
20 days to decisionK013215 · Product code: **BXM** · Anesthesiology
Source: <https://www.510kdatabase.net/k013215/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Stimulator, Nerve, Ac-powered (BXM)
Date received	Sep 26, 2001
Decision date	Oct 16, 2001
Days to decision	20 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Nuvasive, Inc.
Location	San Diego, CA, US
Contact	SEAN M CURRY
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...