FDA 510(k) Premarket Notification

CLEARED

K993239 STRYKER NAVIGATION SYSTEM - NEURO MODULE, MODEL 6000-XXX-XXX

Jan 18, 2000
113 days to decision

K993239 · Product code: **HAW** · Neurology
Source: https://www.510kdatabase.net/k993239/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Neurological Stereotaxic Instrument (HAW)
Date received	Sep 27, 1999
Decision date	Jan 18, 2000
Days to decision	113 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Stryker Corp.**
Location	Mchenry, IL, US
Contact	NICOLE PETTY
510(k) history	124 submissions · 121 cleared · 1976-2023

Stryker Corp. is an American multinational medical technology company headquartered in Portage, Michigan. The company develops and markets surgical equipment, implants, and patient safety technologies used globally across multiple medical specialties. Stryker has received FDA 510(k) clearances from total submissions since its first clearance in 1976. The company maintains active regulatory engagement, with its latest clearance in 2023. Its product portfolio spans orthopedic devices, neurosurgical implants, surgical instruments, and endoscopy systems, reflecting a broad pr...