FDA 510(k) Premarket Notification

**CLEARED**

# K072271 STRYKER NAVIGATION SYSTEM INFINITUS HIP RESURFACING MODULE

Dec 18, 2007
125 days to decision

K072271 · Product code: **HAW** · Neurology
Source: https://www.510kdatabase.net/k072271/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Neurological Stereotaxic Instrument (HAW) |
| Date received | Aug 15, 2007 |
| Decision date | Dec 18, 2007 |
| Days to decision | 125 days |
| Third-party review | No |
| Summary / Statement | Summary |

## APPLICANT

| | |
|---|---|
| Company | **Stryker Corp.** |
| Location | Mchenry, IL, US |
| Contact | BECKY DITTY |
| 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...

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