FDA 510(k) Premarket Notification

# K930224 STRYKER ARTHROSCOPIC ACCESSORIES

**CLEARED**

Jun 30, 1994
531 days to decision

K930224 · Product code: **GCJ** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k930224/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Laparoscope, General & Plastic Surgery (GCJ) |
| Date received | Jan 15, 1993 |
| Decision date | Jun 30, 1994 |
| Days to decision | 531 days |
| Third-party review | No |
| Summary / Statement | Statement |

## APPLICANT

| | |
|---|---|
| Company | **Stryker Corp.** |
| Location | Mchenry, IL, US |
| Contact | SHAUN GOLDEN |
| 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...