

K974771 STRYKER SE5 ARTHROSCOPY SYSTEM WITH HERMES VOICE CONTROLMay 29, 1998
158 days to decisionK974771 · Product code: **GCJ** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k974771/>**SUBMISSION DETAILS**

| | |
|-----------------------|----------------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Laparoscope, General & Plastic Surgery (GCJ) |
| Date received | Dec 22, 1997 |
| Decision date | May 29, 1998 |
| Days to decision | 158 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---------------------------------------------------------------|
| Company | Stryker Endoscopy |
| Location | San Jose, CA, US |
| Contact | SALMAAN HAMEED |
| Website | https://www.stryker.com |
| 510(k) history | 101 submissions · 101 cleared · 1993-2026 |

Stryker Endoscopy is a medical device manufacturer based in San Jose, US. The company specializes in endoscopic and surgical imaging systems for minimally invasive procedures. Stryker Endoscopy has received FDA 510(k) clearances from total submissions since its first clearance in 1993. The company maintains active regulatory status, with its latest clearance in 2026. Its portfolio spans General & Plastic Surgery devices, orthopedic surgical tools, and obstetric and gynecologic instruments. Recent cleared devices include advanced imaging systems such as 4K camera platforms...

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Device record: <https://www.510kdatabase.net/k974771/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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