

K121893 STRYKER SDC3 HD INFORMATION MANAGEMENT SYSTEMSep 5, 2012
68 days to decisionK121893 · Product code: **GCJ** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k121893/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laparoscope, General & Plastic Surgery (GCJ)
Date received	Jun 29, 2012
Decision date	Sep 5, 2012
Days to decision	68 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	Stryker Endoscopy
Location	San Jose, CA, US
Contact	LIFEI LIU
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/k121893/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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