

**K931103 STRYKER STAPLER SYSTEM**Oct 12, 1993  
223 days to decisionK931103 · Product code: **GDW** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k931103/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Staple, Implantable (GDW)
Date received	Mar 3, 1993
Decision date	Oct 12, 1993
Days to decision	223 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Stryker Endoscopy</b>
Location	San Jose, CA, US
Contact	CHARLES L NELSON
Website	<a href="https://www.stryker.com">https://www.stryker.com</a>
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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