

K081995 STRYKER VISION ELECT WHDTV (STRYKER VE WHDTV)Oct 2, 2008
80 days to decisionK081995 · Product code: **GCJ** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k081995/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laparoscope, General & Plastic Surgery (GCJ)
Date received	Jul 14, 2008
Decision date	Oct 2, 2008
Days to decision	80 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Stryker Endoscopy
Location	San Jose, CA, US
Contact	MONICA BARRETT
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/k081995/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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