

**K071859 STRYKER CROSSFIRE SYSTEM**Sep 20, 2007  
77 days to decisionK071859 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k071859/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jul 5, 2007
Decision date	Sep 20, 2007
Days to decision	77 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Stryker Endoscopy</b>
Location	San Jose, CA, US
Contact	K. JEFFREY SEMONE
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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