

K043021 STRYKER URETEROSCOPEMar 16, 2005
133 days to decisionK043021 · Product code: **FGB** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k043021/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ureteroscope And Accessories, Flexible/rigid (FGB)
Date received	Nov 3, 2004
Decision date	Mar 16, 2005
Days to decision	133 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Stryker Endoscopy
Location	San Jose, CA, US
Contact	CHRISTOPHER L COOK
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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