

K061548 STRYKER URETERAL ILLUMINATION SYSTEM IVSep 28, 2006
115 days to decisionK061548 · Product code: **FCS** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k061548/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Catheter, Fiberoptic, Glass, Ureteral (FCS)
Date received	Jun 5, 2006
Decision date	Sep 28, 2006
Days to decision	115 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Stryker Corp.
Location	Mchenry, IL, US
Contact	ERICA A WALTERS
510(k) history	124 submissions · 121 cleared · 1976-2023

Stryker Corp. is an American multinational medical technology company headquartered in Portage, Michigan. The company develops and markets surgical equipment, implants, and patient safety technologies used globally across multiple medical specialties. Stryker has received FDA 510(k) clearances from total submissions since its first clearance in 1976. The company maintains active regulatory engagement, with its latest clearance in 2023. Its product portfolio spans orthopedic devices, neurosurgical implants, surgical instruments, and endoscopy systems, reflecting a broad pr...

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