

K192793 Evis Exera III Colonovideoscope Olympus PCF-H190TL, Evis Exera III Colonovideoscope Olympus PCF-H190TI, Evis Exera III Colonovideoscope Olympus PCF-HQ190L, Evis Exera III Colonovideoscope Olympus PCF-HQ190IJul 17, 2020
291 days to decisionK192793 · Product code: **FDF** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k192793/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Colonoscope And Accessories, Flexible/rigid (FDF)
Date received	Sep 30, 2019
Decision date	Jul 17, 2020
Days to decision	291 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Olympus Medical Systems Corp.
Location	Hachiochi-Shi, JP
Contact	Toshiyuki Nakajima
Website	https://www.olympus-global.com
510(k) history	102 submissions · 102 cleared · 2012-2026

Olympus Medical Systems Corp. is a global medical device manufacturer headquartered in Hachiochi-Shi, Japan. The company specializes in endoscopic imaging systems and therapeutic devices for minimally invasive procedures. Olympus has received FDA 510(k) clearances from total submissions since 2012. The company's regulatory portfolio is dominated by Gastroenterology & Urology devices, including endoscopes, hemostatic forceps, biopsy instruments, and sphincterotomes. The latest clearance in 2026 reflects continued active development and market engagement. Recent cleared dev...

REGULATORY CONSULTANT

Consulting firm	Olympus Corporation of the Americas
Contact	Lisa M Boyle

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)

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Device record: <https://www.510kdatabase.net/k192793/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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