

K252341 EVIS EXERA III BRONCHOVIDEOSCOPE (OLYMPUS BF-Q190)Apr 23, 2026
269 days to decisionK252341 · Product code: **EOQ** · Ear, Nose, Throat
Source: <https://www.510kdatabase.net/k252341/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bronchoscope (flexible Or Rigid) (EOQ)
Date received	Jul 28, 2025
Decision date	Apr 23, 2026
Days to decision	269 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary
Other names	EVIS EXERA III BRONCHOVIDEOSCOPE (OLYMPUS BF-H190); EVIS EXERA III BRONCHOVIDEOSCOPE (OLYMPUS BF-1TH190)

APPLICANT

Company	Olympus Medical Systems Corp.
Location	Hachiochi-Shi, JP
Contact	Shinichiro Kawachi
Website	https://www.olympus-global.com
510(k) history	101 submissions · 101 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	Teffany Hutto

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/k252341/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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