FDA 510(k) Premarket Notification

CLEARED

K251336 VISERA ELITE III Video System Center Olympus OTV-S700 (OTV-S700)

Nov 12, 2025
196 days to decision

K251336 · Product code: **OWN** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k251336/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Confocal Optical Imaging (OWN)
Date received	Apr 30, 2025
Decision date	Nov 12, 2025
Days to decision	196 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary
Other names	VISERA ELITE III Light Source Olympus CLL-S700 (CLL-S700); 4K Camera Head Olympus CH-S700-XZ-EA (CH-S700-XZ-EA)

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 Surgical Technologies of the Americas**
Contact	Darlene Hull

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k251336/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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