

**K022270 MODIFICATION TO OLYMPUS INTEGRATED
ENDOSURGERY SYSTEM ENDOALPHA**Aug 14, 2002
30 days to decisionK022270 · Product code: **ODA** · Radiology
Source: <https://www.510kdatabase.net/k022270/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Endoscopic Central Control Unit (ODA)
Date received	Jul 15, 2002
Decision date	Aug 14, 2002
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Olympus Optical Co., Ltd.
Location	Melville, NY, US
Contact	LAURA STORMS-TYLER
Website	http://www.olympus-global.com/
510(k) history	22 submissions · 22 cleared · 2000-2003

Olympus Optical Co., Ltd. is a global medical device manufacturer headquartered in Melville, US. The company specializes in endoscopic and surgical imaging technologies for minimally invasive procedures. Olympus received FDA 510(k) clearances from total submissions between 2000 and 2003. The company's cleared devices span multiple surgical specialties, with particular strength in endoscopic visualization systems for gastroenterology, urology, otolaryngology, and general surgery. Notable cleared products include bronchofiberscopes, gastrovideoscopes, cystofiberscopes, and ...

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Device record: <https://www.510kdatabase.net/k022270/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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