

**K133538 VISERA CYSTO-NEPHRO VIDEOSCOPE OLYMPUS
 CYF TYPE V2, VISERA CYSTO-NEPHRO VIDEOSCOPE
 OLYMPUS CYF TYPE VA2, VISERA CYSTO-NE**

Aug 7, 2014
 262 days to decision

K133538 · Product code: **NWB** · Gastroenterology & Urology
 Source: <https://www.510kdatabase.net/k133538/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Endoscope, Accessories, Narrow Band Spectrum (NWB)
Date received	Nov 18, 2013
Decision date	Aug 7, 2014
Days to decision	262 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Olympus Corporation of the Americas
Location	Center Valley, PA, US
Contact	SHERI L MUSGNUNG
510(k) history	1 submissions · 1 cleared · 2014-2014

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k133538/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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