

**K252889 Single Use 3-Lumen Needle Knife V (KD-V441M, KD-V451M)**Nov 25, 2025  
76 days to decisionK252889 · Product code: **KNS** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k252889/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Electrosurgical, Endoscopic (with Or Without Accessories) (KNS)
Date received	Sep 10, 2025
Decision date	Nov 25, 2025
Days to decision	76 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Olympus Medical Systems Corporation</b>
Location	Melville, NY, US
Contact	Seiko Yunoki
510(k) history	81 submissions · 81 cleared · 2004-2026

**REGULATORY CONSULTANT**

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Consulting firm	<b>Olympus Corporation of the America</b>
Contact	Jillian Connery

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)510k Database - [www.510kdatabase.net](https://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k252889/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](https://accessdata.fda.gov). - Generated May 13, 2026