

**K223183 Light-Guide Cables**Jan 13, 2023  
93 days to decisionK223183 · Product code: **HBI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k223183/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Illuminator, Fiberoptic, Surgical Field (HBI)
Date received	Oct 12, 2022
Decision date	Jan 13, 2023
Days to decision	93 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Olympus Winter &amp; Ibe GmbH</b>
Location	Melville, NY, US
Contact	Ali Sayad
510(k) history	42 submissions · 42 cleared · 1997-2025

**REGULATORY CONSULTANT**

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Consulting firm	<b>Olympus Surgical Technologies of the Americas</b>
Contact	Christina Flores

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k223183/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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