

**K230350 VISULAS yag**Sep 22, 2023  
225 days to decisionK230350 · Product code: **HQF** · Ophthalmic  
Source: <https://www.510kdatabase.net/k230350/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Ophthalmic (HQF)
Date received	Feb 9, 2023
Decision date	Sep 22, 2023
Days to decision	225 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Carl Zeiss Meditec, Inc.</b>
Location	San Diego, CA, US
Contact	Tanesha Bland
510(k) history	29 submissions · 29 cleared · 1993-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k230350/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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