

**K163632 Katalyst Cyclophotocoagulation Probe**Jun 15, 2017  
175 days to decisionK163632 · Product code: **LQJ** · Ophthalmic  
Source: <https://www.510kdatabase.net/k163632/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lens, Surgical, Laser, Accessory, Ophthalmic Laser (LQJ)
Date received	Dec 22, 2016
Decision date	Jun 15, 2017
Days to decision	175 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Katalyst Surgical, LLC</b>
Location	Washington, DC, US
Contact	Meryl Koch
510(k) history	7 submissions · 7 cleared · 2012-2022

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k163632/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 26, 2026