

**K140362 KATALYST REVOLVER LASER PROBES, KATALYST
REVOLVER ILLUMINATED LASER PROBES, KATALYST
REVOLVER ILLUMINATED PROBES**Sep 15, 2014
215 days to decisionK140362 · Product code: **HQF** · Ophthalmic
Source: <https://www.510kdatabase.net/k140362/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Ophthalmic (HQF)
Date received	Feb 12, 2014
Decision date	Sep 15, 2014
Days to decision	215 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Katalyst Surgical, LLC
Location	Washington, DC, US
Contact	MERYL KOCH
510(k) history	7 submissions · 7 cleared · 2012-2022

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k140362/> Data sourced from FDA 510(k) public records (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. - Generated May 26, 2026