

K241931 OcuMet Beacon (OCUB100)Apr 22, 2025
295 days to decisionK241931 · Product code: **MYC** · Ophthalmic
Source: <https://www.510kdatabase.net/k241931/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ophthalmoscope, Laser, Scanning (MYC)
Date received	Jul 1, 2024
Decision date	Apr 22, 2025
Days to decision	295 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Ocusciences, Inc.
Location	Ann Arbor, MI, US
Contact	Kurt Riegger
510(k) history	1 submissions · 1 cleared · 2025-2025

REGULATORY CONSULTANT

Consulting firm	Healthcare Technologies Consultants, LLC
Contact	Robert Bard

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.netDevice record: <https://www.510kdatabase.net/k241931/> 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 13, 2026