

K242508 Verily Numetric Retinal CameraDec 9, 2024
109 days to decisionK242508 · Product code: **HKI** · Ophthalmic
Source: <https://www.510kdatabase.net/k242508/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Camera, Ophthalmic, Ac-powered (HKI)
Date received	Aug 22, 2024
Decision date	Dec 9, 2024
Days to decision	109 days
Third-party review	Yes
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Verily Life Sciences, LLC
Location	South San Francisco, CA, US
Contact	Shah Pooja
510(k) history	4 submissions · 4 cleared · 2019-2024

REGULATORY CONSULTANT

Consulting firm	Regulatory Technology Services, LLC
Contact	Prithul Bom

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/k242508/> 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