FDA 510(k) Premarket Notification

K242508 Verily Numetric Retinal Camera

CLEARED

Dec 9, 2024
109 days to decision

K242508 · 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