

K243014 EyeQ nanoECPJun 18, 2025
264 days to decisionK243014 · Product code: **HQF** · Ophthalmic
Source: <https://www.510kdatabase.net/k243014/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Ophthalmic (HQF)
Date received	Sep 27, 2024
Decision date	Jun 18, 2025
Days to decision	264 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Statement

APPLICANT

Company	Eyeq, Inc.
Location	Reno, NV, US
Contact	Gary Mocnik
510(k) history	1 submissions · 1 cleared · 2025-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k243014/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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