

K233720 LuminopiaAug 8, 2024
261 days to decisionK233720 · Product code: **QQU** · Ophthalmic
Source: <https://www.510kdatabase.net/k233720/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Digital Therapy Device For Amblyopia (QQU)
Date received	Nov 21, 2023
Decision date	Aug 8, 2024
Days to decision	261 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Luminopia, Inc.
Location	Cambridge, MA, US
Contact	Endri Angjeli
510(k) history	4 submissions · 3 cleared · 2021-2025

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