

K220891 Kahook Dual Blade Glide (KDB Glide)May 17, 2024
781 days to decisionK220891 · Product code: **QUQ** · Ophthalmic
Source: <https://www.510kdatabase.net/k220891/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Knife, Intraocular Pressure Lowering (QUQ)
Date received	Mar 28, 2022
Decision date	May 17, 2024
Days to decision	781 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	New World Medical, Inc.
Location	Rancho Cucamonga, CA, US
Contact	Victor Arellano
510(k) history	12 submissions · 12 cleared · 1993-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k220891/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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