FDA 510(k) Premarket Notification

K220891 Kahook Dual Blade Glide (KDB Glide)

CLEARED

May 17, 2024
781 days to decision

K220891 · 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