

K231051 Ahmed® Glaucoma Valve Model FP8May 12, 2023
29 days to decisionK231051 · Product code: **KYF** · Ophthalmic
Source: <https://www.510kdatabase.net/k231051/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Implant, Eye Valve (KYF)
Date received	Apr 13, 2023
Decision date	May 12, 2023
Days to decision	29 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/k231051/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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