FDA 510(k) Premarket Notification

K093305 ENHANCED ULTRA VIT PROBE

CLEARED

Apr 2, 2010
162 days to decision

K093305 · Product code: **MLZ** · Ophthalmic
Source: https://www.510kdatabase.net/k093305/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Vitrectomy, Instrument Cutter (MLZ)
Date received	Oct 22, 2009
Decision date	Apr 2, 2010
Days to decision	162 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Alcon Research, Ltd.**
Location	Fort Worth, TX, US
Contact	MARTIN A KAUFMAN
510(k) history	16 submissions · 16 cleared · 2000-2017

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k093305/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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