FDA 510(k) Premarket Notification

K170520 HyperVit Vitrectomy Probe, 23 GA, HyperVit Vitrectomy Probe, 25+, HyperVit Vitrectomy Probe, 27+

CLEARED

Mar 22, 2017
28 days to decision

K170520 · Product code: **MLZ** · Ophthalmic
Source: https://www.510kdatabase.net/k170520/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Vitrectomy, Instrument Cutter (MLZ)
Date received	Feb 22, 2017
Decision date	Mar 22, 2017
Days to decision	28 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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