FDA 510(k) Premarket Notification

CLEARED

K000224 FLUROPERM 60-OK, PARAGON HDS-OK

Apr 17, 2000
84 days to decision

K000224 · Product code: **MUW** · Ophthalmic
Source: https://www.510kdatabase.net/k000224/

SUBMISSION DETAILS

Decision	Substantially Equivalent - U
Submission type	Traditional
Device classification	Lens, Contact (orthokeratology) (MUW)
Date received	Jan 24, 2000
Decision date	Apr 17, 2000
Days to decision	84 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Paragon Vision Sciences**
Location	Phoenix, AZ, US
Contact	WILLIAM E MEYERS
510(k) history	15 submissions · 13 cleared · 1994-2018