FDA 510(k) Premarket Notification

K023737 INJECTOR

CLEARED

Jul 14, 2003
249 days to decision

K023737 · Product code: **MSS** · Ophthalmic
Source: https://www.510kdatabase.net/k023737/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Folders And Injectors, Intraocular Lens (iol) (MSS)
Date received	Nov 7, 2002
Decision date	Jul 14, 2003
Days to decision	249 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Anton Meyer & Co. , Ltd.**
Location	Nidau, CH
Contact	THOMAS MEYER
510(k) history	1 submissions · 1 cleared · 2003-2003