FDA 510(k) Premarket Notification

K022560 M2 SINGLE USE MICROKERTOME

CLEARED

Sep 27, 2002
56 days to decision

K022560 · Product code: **HMY** · Ophthalmic
Source: https://www.510kdatabase.net/k022560/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Keratome, Battery-powered (HMY)
Date received	Aug 2, 2002
Decision date	Sep 27, 2002
Days to decision	56 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Moria SA**
Location	Antony, FR
Contact	MELANIE RENAUD-SAMIRI
510(k) history	5 submissions · 5 cleared · 2002-2017