FDA 510(k) Premarket Notification

K020482 KERATOME BLADE 200200 & 600600

CLEARED

Dec 16, 2002
306 days to decision

K020482 · Product code: **HNO** · Ophthalmic
Source: https://www.510kdatabase.net/k020482/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Keratome, Ac-powered (HNO)
Date received	Feb 13, 2002
Decision date	Dec 16, 2002
Days to decision	306 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Micro Specialties, Inc.**
Location	Milford, CT, US
Contact	CHARLES VASSALLO
510(k) history	4 submissions · 4 cleared · 1998-2004