FDA 510(k) Premarket Notification

K980508 KERATOME BLADE, CHIROM MODEL ALK 500500

CLEARED

May 1, 1998
80 days to decision

K980508 · Product code: **HNO** · Ophthalmic
Source: https://www.510kdatabase.net/k980508/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Keratome, Ac-powered (HNO)
Date received	Feb 10, 1998
Decision date	May 1, 1998
Days to decision	80 days
Third-party review	No
Summary / Statement	Summary
Other names	KERATOME BLADE, S.C.M.D. MODEL TUROKERATOME 400400

APPLICANT

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