

K980508 KERATOME BLADE, CHIROM MODEL ALK 500500May 1, 1998
80 days to decisionK980508 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k980508/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 29, 2026