

K942180 KEELER MULTILASE 3000Nov 22, 1994
202 days to decisionK942180 · Product code: **HQF** · Ophthalmic
Source: <https://www.510kdatabase.net/k942180/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Ophthalmic (HQF)
Date received	May 4, 1994
Decision date	Nov 22, 1994
Days to decision	202 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Keeler Instruments, Inc.
Location	Mchenry, IL, US
Contact	EUGENE R VAN ARSDALE
510(k) history	60 submissions · 60 cleared · 1981-2019

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k942180/> 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 20, 2026