

K921414 TROKEL /3 GONIO LASER LENSNov 16, 1992
237 days to decisionK921414 · Product code: **LQJ** · Ophthalmic
Source: <https://www.510kdatabase.net/k921414/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lens, Surgical, Laser, Accessory, Ophthalmic Laser (LQJ)
Date received	Mar 24, 1992
Decision date	Nov 16, 1992
Days to decision	237 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Ocular Instruments, Inc.
Location	Bellevue, WA, US
Contact	TAMSIN J ERICKSON
510(k) history	50 submissions · 50 cleared · 1984-2002

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k921414/> 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