FDA 510(k) Premarket Notification

# K921414 TROKEL /3 GONIO LASER LENS

**CLEARED**

Nov 16, 1992
237 days to decision

K921414 · Product code: **LQJ** · Ophthalmic
Source: https://www.510kdatabase.net/k921414/

## SUBMISSION DETAILS

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|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Lens, Surgical, Laser, Accesssory, 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 |