FDA 510(k) Premarket Notification

# K812219 MODEL 900S INTRAOCULAR PROBE

**CLEARED**

Nov 16, 1981
102 days to decision

K812219 · Product code: **HQF** · Ophthalmic
Source: https://www.510kdatabase.net/k812219/

## SUBMISSION DETAILS

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|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Laser, Ophthalmic (HQF) |
| Date received | Aug 6, 1981 |
| Decision date | Nov 16, 1981 |
| Days to decision | 102 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Lumenis, Inc.** |
| Location | Mchenry, IL, US |
| 510(k) history | 43 submissions · 43 cleared · 1979-2022 |