

K812219 MODEL 900S INTRAOCULAR PROBENov 16, 1981
102 days to decisionK812219 · Product code: **HQF** · Ophthalmic
Source: <https://www.510kdatabase.net/k812219/>**SUBMISSION DETAILS**

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k812219/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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