

K790167 SUNLAMPMar 21, 1979
64 days to decisionK790167 · Product code: **FTC** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k790167/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Ultraviolet, Dermatological (FTC)
Date received	Jan 16, 1979
Decision date	Mar 21, 1979
Days to decision	64 days
Third-party review	No

APPLICANT

Company	Kneilsly Group Ent.
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1979-1979

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Device record: <https://www.510kdatabase.net/k790167/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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