

K803264 SUNLAMPJun 2, 1981
160 days to decisionK803264 · Product code: **FTC** · Radiology
Source: <https://www.510kdatabase.net/k803264/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Ultraviolet, Dermatological (FTC)
Date received	Dec 24, 1980
Decision date	Jun 2, 1981
Days to decision	160 days
Third-party review	No

APPLICANT

Company	Am Labs, Inc.
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1981-1982

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Device record: <https://www.510kdatabase.net/k803264/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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