

K893813 FLOOR STANDING, OPERATING AND EXAMINING LIGHTJul 14, 1989
52 days to decisionK893813 · Product code: **FSS** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k893813/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Surgical, Floor Standing (FSS)
Date received	May 23, 1989
Decision date	Jul 14, 1989
Days to decision	52 days
Third-party review	No

APPLICANT

Company	Astralite Corp.
Location	Diamond Springs, CA, US
Contact	SWARTZ
510(k) history	2 submissions · 2 cleared · 1989-2003

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k893813/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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