FDA 510(k) Premarket Notification

K893813 FLOOR STANDING, OPERATING AND EXAMINING LIGHT

CLEARED

Jul 14, 1989
52 days to decision

K893813 · 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