FDA 510(k) Premarket Notification

K002463 STELLAR

CLEARED

Nov 8, 2000
90 days to decision

K002463 · Product code: **FSY** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k002463/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Surgical, Ceiling Mounted (FSY)
Date received	Aug 10, 2000
Decision date	Nov 8, 2000
Days to decision	90 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Skytron, Div. the Kmw Group, Inc.**
Location	Mchenry, IL, US
Contact	KARI OGREEN
510(k) history	19 submissions · 19 cleared · 1981-2007