

**K002463 STELLAR**Nov 8, 2000  
90 days to decisionK002463 · Product code: **FSY** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k002463/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k002463/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 28, 2026