

**K000639 VISIONARY 2000**May 25, 2000  
90 days to decisionK000639 · Product code: **FSY** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k000639/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Surgical, Ceiling Mounted (FSY)
Date received	Feb 25, 2000
Decision date	May 25, 2000
Days to decision	90 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Burton Medical Products Corp.</b>
Location	Van Nuys, CA, US
Contact	DORIAN SWARTZ
510(k) history	15 submissions · 15 cleared · 1985-2010

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k000639/> 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 20, 2026