

**K893949 #157 EXAM LIGHT**Aug 7, 1989  
67 days to decisionK893949 · Product code: **KFK** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k893949/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Saw, Pneumatically Powered (KFK)
Date received	Jun 1, 1989
Decision date	Aug 7, 1989
Days to decision	67 days
Third-party review	No

**APPLICANT**

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Company	<b>Medmark, Inc.</b>
Location	Versailles, OH, US
Contact	JOHN OLDIGES
510(k) history	3 submissions · 3 cleared · 1989-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k893949/>; 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 30, 2026