

**K896384 FIBEROPTIC LIGHT SOURCE, FOI-1A**Jan 23, 1990  
78 days to decisionK896384 · Product code: **EQH** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k896384/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Source, Carrier, Fiberoptic Light (EQH)
Date received	Nov 6, 1989
Decision date	Jan 23, 1990
Days to decision	78 days
Third-party review	No

**APPLICANT**

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Company	<b>Jedmed Instrument Co.</b>
Location	Mchenry, IL, US
Contact	CRAIG RAPP
510(k) history	92 submissions · 91 cleared · 1979-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k896384/>; 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 May 22, 2026