

**K970878 ENDOLIGHT FIBEROPTIC REGULAR MONOFILAMENT  
AND FINE MONOFILAMENT**May 6, 1997  
57 days to decisionK970878 · Product code: **MPA** · Ophthalmic  
Source: <https://www.510kdatabase.net/k970878/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Endoilluminator (MPA)
Date received	Mar 10, 1997
Decision date	May 6, 1997
Days to decision	57 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>American Medical Devices, Inc.</b>
Location	Atlanta, GA, US
Contact	FRANK J TIGHE
510(k) history	6 submissions · 6 cleared · 1997-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k970878/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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