

**K000728 MEDTRONIC SOLAN FIBEROPTIC CORNEAL LIGHT PROBE**

Mar 21, 2000  
15 days to decision

K000728 · Product code: **FST** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k000728/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Surgical, Fiberoptic (FST)
Date received	Mar 6, 2000
Decision date	Mar 21, 2000
Days to decision	15 days
Third-party review	Yes
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medtronic Xomed</b>
Location	Jacksonville, FL, US
Contact	DAVID TIMLIN
510(k) history	9 submissions · 9 cleared · 1996-2020

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k000728/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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