

**K842609 R.L. MEDICAL TWIN LIGHT SOURCE**Aug 9, 1984  
35 days to decisionK842609 · Product code: **FCW** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k842609/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light Source, Fiberoptic, Routine (FCW)
Date received	Jul 5, 1984
Decision date	Aug 9, 1984
Days to decision	35 days
Third-party review	No

**APPLICANT**

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Company	<b>Tridak Division of Indicon, Inc.</b>
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1984-1990

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

Device record: <https://www.510kdatabase.net/k842609/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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