

**K890436 TECHNOMED PULSOLITH LASER SYSTEM**Mar 23, 1989  
55 days to decisionK890436 · Product code: **FFK** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k890436/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lithotripter, Electro-hydraulic (FFK)
Date received	Jan 27, 1989
Decision date	Mar 23, 1989
Days to decision	55 days
Third-party review	No

**APPLICANT**

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Company	<b>Teknomed, Inc.</b>
Location	Mchenry, IL, US
Contact	RICHARD HUNTER
510(k) history	7 submissions · 7 cleared · 1979-1993

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k890436/>; 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