

**K850298 ULTRALITH ULTRASONIC LITHOTRIPSY SYSTEM**Mar 5, 1985  
40 days to decisionK850298 · Product code: **FEO** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k850298/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lithotripter, Ultrasonic (FEO)
Date received	Jan 24, 1985
Decision date	Mar 5, 1985
Days to decision	40 days
Third-party review	No

**APPLICANT**

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Company	<b>Cabot Medical Corp.</b>
Location	Mchenry, IL, US
Contact	DAVID E GRONOSTAJSK
510(k) history	38 submissions · 38 cleared · 1983-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k850298/>; 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