

**K896822 SURGISMOKE**Jan 11, 1990  
38 days to decisionK896822 · Product code: **FYD** · General Hospital  
Source: <https://www.510kdatabase.net/k896822/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Apparatus, Exhaust, Surgical (FYD)
Date received	Dec 4, 1989
Decision date	Jan 11, 1990
Days to decision	38 days
Third-party review	No

**APPLICANT**

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Company	<b>Spectro Industries, Inc.</b>
Location	Jenkintown, PA, US
Contact	DONALD HEWITT
510(k) history	2 submissions · 2 cleared · 1984-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k896822/>, 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 July 6, 2026