

**K894053 2ND MODIFICATION OF FEF(TM) END-TIDAL CO2  
DETECTOR**Jul 20, 1989  
43 days to decisionK894053 · Product code: **CCK** · Anesthesiology  
Source: <https://www.510kdatabase.net/k894053/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Gas, Carbon-dioxide, Gaseous-phase (CCK)
Date received	Jun 7, 1989
Decision date	Jul 20, 1989
Days to decision	43 days
Third-party review	No

**APPLICANT**

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Company	<b>Fenem, Inc.</b>
Location	New York, NY, US
Contact	SHEILA GLENNON
510(k) history	5 submissions · 5 cleared · 1988-1989

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k894053/> 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 27, 2026