

**K880890 THE S.A.F.E. SYSTEM**May 11, 1988  
70 days to decisionK880890 · Product code: **FYD** · General Hospital  
Source: <https://www.510kdatabase.net/k880890/>**SUBMISSION DETAILS**

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|                       |                                    |
|-----------------------|------------------------------------|
| Decision              | Substantially Equivalent (Cleared) |
| Submission type       | Traditional                        |
| Device classification | Apparatus, Exhaust, Surgical (FYD) |
| Date received         | Mar 2, 1988                        |
| Decision date         | May 11, 1988                       |
| Days to decision      | 70 days                            |
| Third-party review    | No                                 |

**APPLICANT**

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|----------------|-----------------------------------------|
| Company        | <b>Pfizer Laser Systems</b>             |
| Location       | Irvine, CA, US                          |
| Contact        | LINDA DIBENEDETTO                       |
| 510(k) history | 32 submissions · 32 cleared · 1988-1993 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k880890/>, 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 June 30, 2026