

**K920597 HANDI-LASE CO2 SURGICAL LASER INSTRUMENT**Jun 4, 1992  
114 days to decisionK920597 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k920597/>**SUBMISSION DETAILS**

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|-----------------------|---|
| Decision              | Substantially Equivalent (Cleared)      |
| Submission type       | Traditional                             |
| Device classification | Powered Laser Surgical Instrument (GEX) |
| Date received         | Feb 11, 1992                            |
| Decision date         | Jun 4, 1992                             |
| Days to decision      | 114 days                                |
| Third-party review    | No                                      |
| Summary / Statement   | Statement                               |

**APPLICANT**

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|----------------|---------------------------------------|
| Company        | <b>Sorenson Laboratories, Inc.</b>    |
| Location       | Mountain View, CA, US                 |
| Contact        | CHARLES L ROSE                        |
| 510(k) history | 3 submissions · 3 cleared · 1992-1993 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k920597/> 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