

**K131078 SYSTEM 1E LIQUID CHEMICAL STERILANT  
PROCESSING SYSTEM**Jul 5, 2013  
79 days to decisionK131078 · Product code: **MED** · General Hospital  
Source: <https://www.510kdatabase.net/k131078/>**SUBMISSION DETAILS**

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|                       |                                    |
|-----------------------|------------------------------------|
| Decision              | Substantially Equivalent (Cleared) |
| Submission type       | Special                            |
| Device classification | Sterilant, Medical Devices (MED)   |
| Date received         | Apr 17, 2013                       |
| Decision date         | Jul 5, 2013                        |
| Days to decision      | 79 days                            |
| Third-party review    | No                                 |
| Summary / Statement   | Summary                            |

**APPLICANT**

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|----------------|---|
| Company        | <b>STERIS Corporation</b>                 |
| Location       | Mentor, OH, US                            |
| Contact        | MARCIA L BENEDICT                         |
| 510(k) history | 204 submissions · 202 cleared · 1997-2026 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k131078/>; 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 8, 2026