

**K100390 SURGIMAX/ SURGIMAX PLUS**Aug 25, 2010  
190 days to decisionK100390 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k100390/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)                         |
| Submission type       | Traditional  |
| Device classification | Electrosurgical, Cutting & Coagulation & Accessories (GEI) |
| Date received         | Feb 16, 2010   |
| Decision date         | Aug 25, 2010   |
| Days to decision      | 190 days   |
| Third-party review    | No   |
| Summary / Statement   | Summary  |

**APPLICANT**

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|----------------|---------------------------------------|
| Company        | <b>Elliquence, LLC</b>                |
| Location       | Shelton, CT, US                       |
| Contact        | JOSEPH AZARY                          |
| 510(k) history | 8 submissions · 8 cleared · 2010-2025 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k100390/> 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