

**K170107 Surgi Max Ultra**May 3, 2017  
111 days to decisionK170107 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k170107/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jan 12, 2017
Decision date	May 3, 2017
Days to decision	111 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Elliquence, LLC</b>
Location	Shelton, CT, US
Contact	Paul D. Buhrke IV
510(k) history	8 submissions · 8 cleared · 2010-2025

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k170107/> 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