

**K001253 SURGITRON IEC II**May 26, 2000  
37 days to decisionK001253 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k001253/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Apr 19, 2000
Decision date	May 26, 2000
Days to decision	37 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Ellman Intl., Inc.</b>
Location	Hewlett, NY, US
Contact	FRANK LIN
510(k) history	15 submissions · 15 cleared · 1996-2001

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k001253/> 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 29, 2026