

**K120510 TRANSCEND**Jul 19, 2013  
514 days to decisionK120510 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k120510/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Feb 21, 2012
Decision date	Jul 19, 2013
Days to decision	514 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Syneron Medical, Ltd.</b>
Location	Yokneam Elite, IL
Contact	SAM WADE
510(k) history	35 submissions · 35 cleared · 2002-2017

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k120510/> 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 1, 2026