

**K022266 AURORA SR**Oct 3, 2002  
83 days to decisionK022266 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k022266/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Jul 12, 2002
Decision date	Oct 3, 2002
Days to decision	83 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Syneron Medical, Ltd.</b>
Location	Yokneam Elite, IL
Contact	AMIR WALDMAN
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/k022266/> 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