

**K032514 AC APPLICATOR, AURORA AC**Jan 27, 2004  
166 days to decisionK032514 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k032514/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Aug 14, 2003
Decision date	Jan 27, 2004
Days to decision	166 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k032514/> 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