

**K070092 MODIFICATION TO VELASMOOTH, SHAPER**Jul 27, 2007  
198 days to decisionK070092 · Product code: **NUV** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k070092/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Massager, Vacuum, Light Induced Heating (NUV)
Date received	Jan 10, 2007
Decision date	Jul 27, 2007
Days to decision	198 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Syneron Medical, Ltd.</b>
Location	Yokneam Elite, IL
Contact	YONI IGER
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/k070092/> 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 28, 2026