

**K061720 COSMELIGHT**Jan 31, 2007  
226 days to decisionK061720 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k061720/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Jun 19, 2006
Decision date	Jan 31, 2007
Days to decision	226 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Penntack Enterprises, Inc.</b>
Location	Miami, FL, US
Contact	PABLO PENA
510(k) history	1 submissions · 1 cleared · 2007-2007

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k061720/> 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