

**K070466 REVOLIX DUO LASER SYSTEM**Apr 17, 2007  
60 days to decisionK070466 · Product code: **GEX** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k070466/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Feb 16, 2007
Decision date	Apr 17, 2007
Days to decision	60 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Allmed Systems, Inc.</b>
Location	Pleasanton, CA, US
Contact	PETER ALLEN
510(k) history	8 submissions · 8 cleared · 2004-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k070466/> 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 7, 2026