

**K063072 CANDELA FAMILY OF Q-SWITCHED ALEXANDRITE  
LASER (ALEXLAZR) SYSTEMS: 532 AND 1064 ND:YAG  
HANDPIECE ACCESSORIES**Dec 20, 2006  
75 days to decisionK063072 · Product code: **GEX** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k063072/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Oct 6, 2006
Decision date	Dec 20, 2006
Days to decision	75 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Candela Corp.</b>
Location	Natick, MA, US
Contact	JEFFERY ROBERTS
510(k) history	48 submissions · 48 cleared · 1986-2019

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k063072/> 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 18, 2026