

K081324 CANDELA FAMILY OF Q-SWITCHED ALEXANDRITE LASERSJun 17, 2008
36 days to decisionK081324 · Product code: **GEX** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k081324/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	May 12, 2008
Decision date	Jun 17, 2008
Days to decision	36 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Candela Corp.
Location	Natick, MA, US
Contact	LISA LACROIX
510(k) history	48 submissions · 48 cleared · 1986-2019

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k081324/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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