

K043251 CANDELA FAMILY OF PULSED DYE LASER SYSTEMS, MODELS VBEAM, CBEAM, SPTL-1BFeb 9, 2005
78 days to decisionK043251 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k043251/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Nov 23, 2004
Decision date	Feb 9, 2005
Days to decision	78 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k043251/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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