

**K212073 Alma Diode Tabletop Laser**Jul 14, 2022  
377 days to decisionK212073 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k212073/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Jul 2, 2021
Decision date	Jul 14, 2022
Days to decision	377 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Alma Lasers , Ltd.</b>
Location	Pleasanton, CA, US
Contact	Avi Hirshnzon
510(k) history	14 submissions · 14 cleared · 2007-2022

**REGULATORY CONSULTANT**

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Consulting firm	<b>Hoy &amp; Associates Regulatory Consulting</b>
Contact	Connie Hoy

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov

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