

**K932065 DLASE 300 UPGRADE**Jul 22, 1993  
84 days to decisionK932065 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k932065/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Apr 29, 1993
Decision date	Jul 22, 1993
Days to decision	84 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Incirt Tech., Inc.</b>
Location	San Carlos, CA, US
Contact	MICHAEL YESSIK
510(k) history	1 submissions · 1 cleared · 1993-1993

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k932065/> 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 July 6, 2026