

**K944626 MODEL 210-AS ASPIRATING  
ENDOPHOTOCOAGULATOR LASER**Nov 3, 1994  
44 days to decisionK944626 · Product code: **GEX** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k944626/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Sep 20, 1994
Decision date	Nov 3, 1994
Days to decision	44 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Mnlase, Inc.</b>
Location	Minneapolis, MN, US
Contact	BRUCE HEYMANN
510(k) history	22 submissions · 22 cleared · 1992-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k944626/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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