

**K082721 LASERPRO 810, 940 AND 980 DIODE LASER SYSTEMS**Oct 28, 2008  
41 days to decisionK082721 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k082721/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Sep 17, 2008
Decision date	Oct 28, 2008
Days to decision	41 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Photo Medex, Inc.</b>
Location	Carlsbad, CA, US
Contact	ALFRED INTINTOLI
510(k) history	10 submissions · 10 cleared · 2001-2008

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k082721/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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