

**K063384 DIODENT MICRO DENTAL LASER SYSTEM, MODELS 810 AND 980**

Mar 16, 2007  
128 days to decision

K063384 · Product code: **GEX** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k063384/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Nov 8, 2006
Decision date	Mar 16, 2007
Days to decision	128 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Hoya Conbio, Inc.</b>
Location	Fremont, CA, US
Contact	LIZA BURNS
510(k) history	7 submissions · 7 cleared · 2006-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k063384/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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