FDA 510(k) Premarket Notification

K073332 V-RASER DIODE LASER SYSTEM

CLEARED

Feb 14, 2008
79 days to decision

K073332 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k073332/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Nov 27, 2007
Decision date	Feb 14, 2008
Days to decision	79 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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