FDA 510(k) Premarket Notification

K082586 LIGHTWAVE PROFESSIONAL DELUXE

CLEARED

Jan 4, 2010
483 days to decision

K082586 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k082586/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Sep 8, 2008
Decision date	Jan 4, 2010
Days to decision	483 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Lightwave Technologies, LLC**
Location	Great Neck, NY, US
Contact	MARIA F GRIFFIN
510(k) history	1 submissions · 1 cleared · 2010-2010

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k082586/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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