

K180888 LYFTAug 30, 2018
148 days to decisionK180888 · Product code: **OLP** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k180888/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Over-the-counter Powered Light Based Laser For Acne (OLP)
Date received	Apr 4, 2018
Decision date	Aug 30, 2018
Days to decision	148 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Uplevity, Inc.
Location	Niantic, CT, US
Contact	Thomas Kraft
510(k) history	1 submissions · 1 cleared · 2018-2018

REGULATORY CONSULTANT

Consulting firm	Contract In-House Counsel and Consultants, LLC
Contact	Marc C. Sanchez

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k180888/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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