

**K180900 LED Light Therapy Device**Jul 10, 2018  
95 days to decisionK180900 · Product code: **OLP** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k180900/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Over-the-counter Powered Light Based Laser For Acne (OLP)
Date received	Apr 6, 2018
Decision date	Jul 10, 2018
Days to decision	95 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Uvbiotek, LLC</b>
Location	Hudson, NY, US
Contact	Dave Oberhelman
510(k) history	2 submissions · 2 cleared · 2017-2018

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k180900/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 30, 2026