

K252960 SOLIUS PRO UVB Light PanelJan 16, 2026
122 days to decisionK252960 · Product code: **SGZ** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k252960/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ultraviolet Lamp Products (SGZ)
Date received	Sep 16, 2025
Decision date	Jan 16, 2026
Days to decision	122 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Solius Labs, Inc.
Location	Bainbridge Island, WA, US
Contact	Chris Kiple
510(k) history	1 submissions · 1 cleared · 2026-2026

REGULATORY CONSULTANT

Consulting firm	Bentley Biomedical Consulting, LLC
Contact	Shepard Bentley

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/k252960/> Data sourced from FDA 510(k) public records (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. - Generated May 13, 2026