

K253683 FAQ™ (LED Panel)Feb 20, 2026
91 days to decisionK253683 · Product code: **OHS** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k253683/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light Based Over The Counter Wrinkle Reduction (OHS)
Date received	Nov 21, 2025
Decision date	Feb 20, 2026
Days to decision	91 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Statement
Other names	FAQ™ (Dual LED Panel)

APPLICANT

Company	Foreo, Inc.
Location	Las Vegas, NV, US
Contact	Feldstein Evan
510(k) history	10 submissions · 10 cleared · 2016-2026

REGULATORY CONSULTANT

Consulting firm	DD Consulting
Contact	Domljanovic Danijela

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

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