

K243587 PureLift GLOWMar 6, 2025
106 days to decisionK243587 · Product code: **NFO** · Neurology
Source: <https://www.510kdatabase.net/k243587/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Transcutaneous Electrical, Aesthetic Purposes (NFO)
Date received	Nov 20, 2024
Decision date	Mar 6, 2025
Days to decision	106 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Xtreem Pulse, LLC
Location	New York, NY, US
Contact	Andrew Barile
510(k) history	4 submissions · 4 cleared · 2018-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k243587/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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