

**K221443 PureLift Pro Plus**Oct 21, 2022  
156 days to decisionK221443 · Product code: **NFO** · Neurology  
Source: <https://www.510kdatabase.net/k221443/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Stimulator, Transcutaneous Electrical, Aesthetic Purposes (NFO)
Date received	May 18, 2022
Decision date	Oct 21, 2022
Days to decision	156 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k221443/> 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 26, 2026