

**K212238 TheraFace Microcurrent**Sep 29, 2021  
72 days to decisionK212238 · Product code: **NFO** · Neurology  
Source: <https://www.510kdatabase.net/k212238/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Transcutaneous Electrical, Aesthetic Purposes (NFO)
Date received	Jul 19, 2021
Decision date	Sep 29, 2021
Days to decision	72 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Theragun, Inc.</b>
Location	Los Angeles, CA, US
Contact	Jaime Sanchez
510(k) history	3 submissions · 3 cleared · 2021-2021

**REGULATORY CONSULTANT**

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Consulting firm	<b>Schiff &amp; Company, Inc.</b>
Contact	Thomas Padula

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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