

**K232001 Nu Skin RenuSpa iO**Nov 21, 2023  
139 days to decisionK232001 · Product code: **NFO** · Neurology  
Source: <https://www.510kdatabase.net/k232001/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Transcutaneous Electrical, Aesthetic Purposes (NFO)
Date received	Jul 5, 2023
Decision date	Nov 21, 2023
Days to decision	139 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Nse Products, Inc.</b>
Location	Provo, UT, US
Contact	McKay Noll
510(k) history	1 submissions · 1 cleared · 2023-2023

**REGULATORY CONSULTANT**

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Consulting firm	<b>Methodize, Inc.</b>
Contact	Jack Slovick

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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