

**K222386 Visionbody**Feb 24, 2023  
200 days to decisionK222386 · Product code: **NGX** · Physical MedicineSource: <https://www.510kdatabase.net/k222386/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Muscle, Powered, For Muscle Conditioning (NGX)
Date received	Aug 8, 2022
Decision date	Feb 24, 2023
Days to decision	200 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Vb Technologies AG</b>
Location	Zug, CH
Contact	Jorge Millan
510(k) history	1 submissions · 1 cleared · 2023-2023

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k222386/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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