

K151903 Slendertone connect AbsNov 6, 2015
119 days to decisionK151903 · Product code: **NGX** · Physical MedicineSource: <https://www.510kdatabase.net/k151903/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Muscle, Powered, For Muscle Conditioning (NGX)
Date received	Jul 10, 2015
Decision date	Nov 6, 2015
Days to decision	119 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Bio-Medical Research, Ltd.
Location	Washington Dc, DC, US
Contact	ANNE-MARIE KEENAN
510(k) history	32 submissions · 31 cleared · 1996-2021

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k151903/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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