

K100556 SLENDERTONE FLEX MAX, MODEL 517 USAug 2, 2010
154 days to decisionK100556 · Product code: **NGX** · Physical MedicineSource: <https://www.510kdatabase.net/k100556/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Muscle, Powered, For Muscle Conditioning (NGX)
Date received	Mar 1, 2010
Decision date	Aug 2, 2010
Days to decision	154 days
Third-party review	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/k100556/> Data sourced from FDA 510(k) public records (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. - Generated June 15, 2026