

K151722 Slim UP ULTRAMar 18, 2016
267 days to decisionK151722 · Product code: **NGX** · Physical MedicineSource: <https://www.510kdatabase.net/k151722/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Muscle, Powered, For Muscle Conditioning (NGX)
Date received	Jun 25, 2015
Decision date	Mar 18, 2016
Days to decision	267 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Aime Medical, Inc.
Location	Largo, FL, US
Contact	PAUL GUILBAUD
510(k) history	1 submissions · 1 cleared · 2016-2016

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k151722/> 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 3, 2026