

K171035 Katalyst Mark 1 Muscle Stimulation SystemSep 6, 2017
153 days to decisionK171035 · Product code: **NGX** · Physical Medicine
Source: <https://www.510kdatabase.net/k171035/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Muscle, Powered, For Muscle Conditioning (NGX)
Date received	Apr 6, 2017
Decision date	Sep 6, 2017
Days to decision	153 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Katalyst, Inc.
Location	Seattle, WA, US
Contact	Amish Patel
510(k) history	3 submissions · 3 cleared · 2017-2020

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k171035/> 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 28, 2026