

K173682 L300 Go SystemMar 9, 2018
98 days to decisionK173682 · Product code: **GZI** · Neurology
Source: <https://www.510kdatabase.net/k173682/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Neuromuscular, External Functional (GZI)
Date received	Dec 1, 2017
Decision date	Mar 9, 2018
Days to decision	98 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Bioness, Inc.
Location	Valencia, CA, US
Contact	Mercedes Bayani
510(k) history	13 submissions · 13 cleared · 2011-2022

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k173682/> 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 May 27, 2026