

**K230701 Stim2Go**Nov 24, 2023  
255 days to decisionK230701 · Product code: **IPF** · Physical MedicineSource: <https://www.510kdatabase.net/k230701/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Muscle, Powered (IPF)
Date received	Mar 14, 2023
Decision date	Nov 24, 2023
Days to decision	255 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>PAJUNK GmbH Medizintechnologie</b>
Location	Geisingen, DE
Contact	Christian Quass
510(k) history	44 submissions · 44 cleared · 2006-2025

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k230701/> 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 14, 2026