

K223802 accufitOct 6, 2023
291 days to decisionK223802 · Product code: **IPF** · Physical MedicineSource: <https://www.510kdatabase.net/k223802/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Muscle, Powered (IPF)
Date received	Dec 19, 2022
Decision date	Oct 6, 2023
Days to decision	291 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Mettler Electronics Corporation
Location	Anaheim, CA, US
Contact	An Le
510(k) history	2 submissions · 2 cleared · 2023-2024

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k223802/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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