

K231032 AMP by Method Mobility, AMP Plus (or +) by Method MobilityMay 10, 2023
29 days to decisionK231032 · Product code: ITI · Physical Medicine
Source: <https://www.510kdatabase.net/k231032/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wheelchair, Powered (ITI)
Date received	Apr 11, 2023
Decision date	May 10, 2023
Days to decision	29 days
Third-party review	Yes
Combination product	No
PCCP authorized	No
Summary / Statement	Statement

APPLICANT

Company	Method Mobility
Location	Fresno, CA, US
Contact	Paul Dickie
510(k) history	1 submissions · 1 cleared · 2023-2023

REGULATORY CONSULTANT

Consulting firm	Third Party Review Group, LLC
Contact	Dave Yungvirt

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k231032/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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