

K200583 Ki Mobility Focus CR, Ki Mobility Focus CRe, Ki Mobility Focus CR TTL

Sep 5, 2023
1278 days to decision

K200583 · Product code: **IOR** · Physical Medicine
Source: <https://www.510kdatabase.net/k200583/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wheelchair, Mechanical (IOR)
Date received	Mar 6, 2020
Decision date	Sep 5, 2023
Days to decision	1278 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Ki Mobility, LLC
Location	Stevens Point, WI, US
Contact	Mark Murphy
510(k) history	5 submissions · 5 cleared · 2006-2023

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k200583/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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