FDA 510(k) Premarket Notification

K210920 iBOT Personal Mobility Device (iBOT PMD)

CLEARED

Jun 16, 2021
79 days to decision

K210920 · Product code: **IMK** · Physical Medicine
Source: https://www.510kdatabase.net/k210920/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wheelchair, Stair Climbing (IMK)
Date received	Mar 29, 2021
Decision date	Jun 16, 2021
Days to decision	79 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Mobius Mobility**
Location	Manchester, NH, US
Contact	Joseph Sullivan
510(k) history	2 submissions · 2 cleared · 2021-2025

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k210920/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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