FDA 510(k) Premarket Notification

CLEARED

K220740 Power Wheelchair, W5907 (Q50 R Carbon)

May 12, 2022
59 days to decision

K220740 · Product code: **ITI** · Physical Medicine
Source: https://www.510kdatabase.net/k220740/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wheelchair, Powered (ITI)
Date received	Mar 14, 2022
Decision date	May 12, 2022
Days to decision	59 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Zhejiang Innuovo Rehabilitation Devices Co.,Ltd**
Location	Dongyang, CN
Contact	Leo Zheng
510(k) history	16 submissions · 16 cleared · 2022-2026

REGULATORY CONSULTANT

Consulting firm	**Shanghai Sungo Management Consulting Company Limited.**
Contact	Ivy Wang

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