

K203552 Rapid RebootDec 22, 2020
18 days to decisionK203552 · Product code: **IRP** · Physical Medicine
Source: <https://www.510kdatabase.net/k203552/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Massager, Powered Inflatable Tube (IRP)
Date received	Dec 4, 2020
Decision date	Dec 22, 2020
Days to decision	18 days
Third-party review	Yes
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Rapid Reboot Recovery Products, LLC
Location	Lindon, UT, US
Contact	David Johnson
510(k) history	2 submissions · 2 cleared · 2018-2020

REGULATORY CONSULTANT

Consulting firm	Regulatory Technology Services, LLC
Contact	Prithul Bom

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

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