

**K211745 RecoveryAir PRO**Oct 4, 2021  
119 days to decisionK211745 · Product code: **IRP** · Physical Medicine  
Source: <https://www.510kdatabase.net/k211745/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Massager, Powered Inflatable Tube (IRP)
Date received	Jun 7, 2021
Decision date	Oct 4, 2021
Days to decision	119 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Theragun, Inc.</b>
Location	Los Angeles, CA, US
Contact	Jaime Sanchez
510(k) history	3 submissions · 3 cleared · 2021-2021

**REGULATORY CONSULTANT**

---

Consulting firm	<b>Schiff &amp; Company, Inc.</b>
Contact	Thomas Padula

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k211745/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](https://accessdata.fda.gov). - Generated May 24, 2026