

**K210979 Prelivia**May 13, 2021  
42 days to decisionK210979 · Product code: **IPF** · Physical MedicineSource: <https://www.510kdatabase.net/k210979/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Muscle, Powered (IPF)
Date received	Apr 1, 2021
Decision date	May 13, 2021
Days to decision	42 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Rehabtronics, Inc.</b>
Location	Edmonton, CA
Contact	Mike Pearson
510(k) history	2 submissions · 2 cleared · 2016-2021

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k210979/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 27, 2026