

**K173530 Indego(R)**Jan 31, 2018  
77 days to decisionK173530 · Product code: **PHL** · Neurology  
Source: <https://www.510kdatabase.net/k173530/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Exoskeleton (PHL)
Date received	Nov 15, 2017
Decision date	Jan 31, 2018
Days to decision	77 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Parker Hannifin Corporation</b>
Location	Macedonia, OH, US
Contact	Achilleas Dorotheou
510(k) history	5 submissions · 5 cleared · 2016-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k173530/> 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 25, 2026