

**K221859 Atalante**Dec 29, 2022  
185 days to decisionK221859 · Product code: **PHL** · Neurology  
Source: <https://www.510kdatabase.net/k221859/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Exoskeleton (PHL)
Date received	Jun 27, 2022
Decision date	Dec 29, 2022
Days to decision	185 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Wandercraft SAS</b>
Location	Paris, FR
Contact	Matthieu Masselin
510(k) history	3 submissions · 3 cleared · 2022-2025

**REGULATORY CONSULTANT**

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Consulting firm	<b>Emergo Global Consulting, LLC</b>
Contact	Andre Kindsvater

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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