

**K170564 MyndMove System, MyndMove Functional Electrical Stimulator**Aug 30, 2017  
184 days to decisionK170564 · Product code: **GZI** · Neurology  
Source: <https://www.510kdatabase.net/k170564/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Neuromuscular, External Functional (GZI)
Date received	Feb 27, 2017
Decision date	Aug 30, 2017
Days to decision	184 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Myndtec, Inc.</b>
Location	Mississauga, CA
Contact	Alexa Granger
510(k) history	3 submissions · 3 cleared · 2017-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k170564/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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