

**K132416 REHASTIM 2, ERIGOPRO (FES)**May 28, 2014  
299 days to decisionK132416 · Product code: **GZI** · Neurology  
Source: <https://www.510kdatabase.net/k132416/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Neuromuscular, External Functional (GZI)
Date received	Aug 2, 2013
Decision date	May 28, 2014
Days to decision	299 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Hasomed GmbH</b>
Location	Magdeburg, DE
Contact	MATTHIAS WEBER
510(k) history	4 submissions · 4 cleared · 2007-2017

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k132416/> 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 June 1, 2026