

**K123354 REVITIVE IX**Jun 28, 2013  
240 days to decisionK123354 · Product code: **IPF** · Physical Medicine  
Source: <https://www.510kdatabase.net/k123354/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Muscle, Powered (IPF)
Date received	Oct 31, 2012
Decision date	Jun 28, 2013
Days to decision	240 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Actegy , Ltd.</b>
Location	Washington, DC, US
Contact	JOHN J SMITH
510(k) history	7 submissions · 7 cleared · 2013-2021

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k123354/> 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 26, 2026