

**K133453 IOVERA**Mar 24, 2014  
132 days to decisionK133453 · Product code: **GXH** · Neurology  
Source: <https://www.510kdatabase.net/k133453/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Surgical, Cryogenic (GXH)
Date received	Nov 12, 2013
Decision date	Mar 24, 2014
Days to decision	132 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Myoscience, Inc.</b>
Location	Redwood City, CA, US
Contact	BIJESH CHANDRAN
510(k) history	9 submissions · 9 cleared · 2009-2018

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k133453/> 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 31, 2026