

**K102610 FOCUS EMG DEVICE**Mar 4, 2011  
175 days to decisionK102610 · Product code: **IKN** · Neurology  
Source: <https://www.510kdatabase.net/k102610/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electromyograph, Diagnostic (IKN)
Date received	Sep 10, 2010
Decision date	Mar 4, 2011
Days to decision	175 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Teleemg, LLC USA</b>
Location	Woburn, MA, US
Contact	JOE F JABRE, M.D.
510(k) history	2 submissions · 2 cleared · 2011-2015

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k102610/> 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 5, 2026