

**K142624 Neuromaster G1 MEE200**Apr 24, 2015  
220 days to decisionK142624 · Product code: **GWF** · Neurology  
Source: <https://www.510kdatabase.net/k142624/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Electrical, Evoked Response (GWF)
Date received	Sep 16, 2014
Decision date	Apr 24, 2015
Days to decision	220 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Nihon Kohden Corporation</b>
Location	Tokyo, JP
Contact	Natalie Kennel
510(k) history	18 submissions · 18 cleared · 2015-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k142624/> 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 24, 2026