

**K031930 HIGH SPEED EMG MODULE**Mar 3, 2004  
254 days to decisionK031930 · Product code: **GWF** · Neurology  
Source: <https://www.510kdatabase.net/k031930/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Electrical, Evoked Response (GWF)
Date received	Jun 23, 2003
Decision date	Mar 3, 2004
Days to decision	254 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Medical Measurement Systems B.V.</b>
Location	Enschede Overijssel, NL
Contact	KARIN OGINK-SOMHORST
510(k) history	7 submissions · 7 cleared · 2004-2012

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k031930/> 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 18, 2026