

**K871338 NOVAMETRIX MODEL 200 MAGSTIM**Aug 4, 1987  
123 days to decisionK871338 · Product code: **GWF** · Neurology  
Source: <https://www.510kdatabase.net/k871338/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Electrical, Evoked Response (GWF)
Date received	Apr 3, 1987
Decision date	Aug 4, 1987
Days to decision	123 days
Third-party review	No

**APPLICANT**

---

Company	<b>Novamatrix Medical Systems, Inc.</b>
Location	Mchenry, IL, US
Contact	JAMES R VEALE
510(k) history	45 submissions · 45 cleared · 1978-2001

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k871338/>; 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 25, 2026