

**K860903 TM-3600 A NEURODIAGNOSTIC SYSTEM**Jun 5, 1986  
87 days to decisionK860903 · Product code: **GWF** · Neurology  
Source: <https://www.510kdatabase.net/k860903/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Electrical, Evoked Response (GWF)
Date received	Mar 10, 1986
Decision date	Jun 5, 1986
Days to decision	87 days
Third-party review	No

**APPLICANT**

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Company	<b>Tracor Northern, Inc.</b>
Location	Mchenry, IL, US
Contact	PATTY SONNTAG
510(k) history	10 submissions · 10 cleared · 1981-1988

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k860903/>; 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 July 1, 2026