

**K870567 EVOPORT 300**Oct 6, 1987  
238 days to decisionK870567 · Product code: **GWF** · Neurology  
Source: <https://www.510kdatabase.net/k870567/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Electrical, Evoked Response (GWF)
Date received	Feb 10, 1987
Decision date	Oct 6, 1987
Days to decision	238 days
Third-party review	No

**APPLICANT**

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Company	<b>Dantec Electronics, Inc.</b>
Location	Walker, MI, US
Contact	HENRIK HENRIKSEN
510(k) history	12 submissions · 12 cleared · 1985-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k870567/>; 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