

**K093304 PROTEKTOR 32**Aug 11, 2010  
294 days to decisionK093304 · Product code: **GWF** · Neurology  
Source: <https://www.510kdatabase.net/k093304/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Electrical, Evoked Response (GWF)
Date received	Oct 21, 2009
Decision date	Aug 11, 2010
Days to decision	294 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Natus Medical, Inc.</b>
Location	San Carlos, CA, US
Contact	GOLDY SINGH
510(k) history	12 submissions · 12 cleared · 1995-2010

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k093304/> 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 June 20, 2026