

**K023525 RETISCAN, RETIPORT**Mar 26, 2004  
522 days to decisionK023525 · Product code: **GWE** · Neurology  
Source: <https://www.510kdatabase.net/k023525/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Stimulator, Photic, Evoked Response (GWE)
Date received	Oct 21, 2002
Decision date	Mar 26, 2004
Days to decision	522 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Roland Consult</b>
Location	Brandenburg, DE
Contact	MATTHIAS MAI
510(k) history	1 submissions · 1 cleared · 2004-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k023525/> 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 30, 2026