

**K122149 QBTES**Oct 17, 2012  
90 days to decisionK122149 · Product code: **LQD** · Neurology  
Source: <https://www.510kdatabase.net/k122149/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Recorder, Attention Task Performance (LQD)
Date received	Jul 19, 2012
Decision date	Oct 17, 2012
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Qbtech AB</b>
Location	Goteborg, SE
Contact	HANS BOSTROM
510(k) history	4 submissions · 4 cleared · 2004-2016

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k122149/> 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 3, 2026