

**K093092 ODYSSEY WORKSTATION**Jan 21, 2010  
112 days to decisionK093092 · Product code: **DQK** · CardiovascularSource: <https://www.510kdatabase.net/k093092/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Computer, Diagnostic, Programmable (DQK)
Date received	Oct 1, 2009
Decision date	Jan 21, 2010
Days to decision	112 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Stereotaxis, Inc.</b>
Location	St. Louis, MO, US
Contact	DENNIS POZZO
510(k) history	28 submissions · 28 cleared · 2002-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k093092/> 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 May 14, 2026