

**K150782 AbStats Gateway**Dec 14, 2015  
264 days to decisionK150782 · Product code: **DQD** · Cardiovascular  
Source: <https://www.510kdatabase.net/k150782/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stethoscope, Electronic (DQD)
Date received	Mar 25, 2015
Decision date	Dec 14, 2015
Days to decision	264 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Gi Logic, Inc.</b>
Location	Pasadena, CA, US
Contact	James Beeton
510(k) history	1 submissions · 1 cleared · 2015-2015

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k150782/> 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