

**K231551 Stethophone**Oct 12, 2023  
135 days to decisionK231551 · Product code: **DQD** · Cardiovascular  
Source: <https://www.510kdatabase.net/k231551/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stethoscope, Electronic (DQD)
Date received	May 30, 2023
Decision date	Oct 12, 2023
Days to decision	135 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Sparrow Acoustics, Inc.</b>
Location	Lucasville, CA
Contact	Nadia Ivanova
510(k) history	4 submissions · 4 cleared · 2023-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k231551/> 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 25, 2026