

**K082771 PROTRON TECHNOLOGIES, LLC STETHOTRON**Feb 27, 2009  
158 days to decisionK082771 · Product code: **DQD** · CardiovascularSource: <https://www.510kdatabase.net/k082771/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stethoscope, Electronic (DQD)
Date received	Sep 22, 2008
Decision date	Feb 27, 2009
Days to decision	158 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Protron Technologies, LLC</b>
Location	Leonia, NJ, US
Contact	LEON M DONDYSH
510(k) history	1 submissions · 1 cleared · 2009-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k082771/> 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 23, 2026