

**K192607 LiteWalk**Dec 19, 2019  
90 days to decisionK192607 · Product code: **IRP** · Physical Medicine  
Source: <https://www.510kdatabase.net/k192607/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Massager, Powered Inflatable Tube (IRP)
Date received	Sep 20, 2019
Decision date	Dec 19, 2019
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Viasonix , Ltd.</b>
Location	Yonkers, NY, US
Contact	Shlomi Deler
510(k) history	6 submissions · 6 cleared · 2011-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k192607/> 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 26, 2026