

**K161919 UltraScan 650**Apr 5, 2017  
266 days to decisionK161919 · Product code: **MUA** · Radiology  
Source: <https://www.510kdatabase.net/k161919/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bone Sonometer (MUA)
Date received	Jul 13, 2016
Decision date	Apr 5, 2017
Days to decision	266 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Cyberlogic, Inc.</b>
Location	New York, NY, US
Contact	Jonathan J. Kaufman
510(k) history	1 submissions · 1 cleared · 2017-2017

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k161919/> 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 16, 2026