

**K173265 UltraVision 2 Diagnostic Ultrasound System**Nov 7, 2017  
27 days to decisionK173265 · Product code: IYN · Radiology  
Source: <https://www.510kdatabase.net/k173265/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	System, Imaging, Pulsed Doppler, Ultrasonic (IYN)
Date received	Oct 11, 2017
Decision date	Nov 7, 2017
Days to decision	27 days
Third-party review	Yes
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Winprobe Corporation</b>
Location	Palm Beach Gardens, FL, US
Contact	Guy Scott
510(k) history	2 submissions · 2 cleared · 2015-2017

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k173265/> 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