

**K102813 EZONO 3000**Jun 30, 2011  
275 days to decisionK102813 · Product code: **IYN** · Radiology  
Source: <https://www.510kdatabase.net/k102813/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Doppler, Ultrasonic (IYN)
Date received	Sep 28, 2010
Decision date	Jun 30, 2011
Days to decision	275 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Life-Tech, Inc.</b>
Location	Stafford, TX, US
Contact	SKIP RIMER
510(k) history	14 submissions · 14 cleared · 2005-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k102813/> 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 17, 2026