

**K102637 DYNAREX ULTRASOUND GEL**Feb 16, 2011  
155 days to decisionK102637 · Product code: **MUI** · Radiology  
Source: <https://www.510kdatabase.net/k102637/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Media, Coupling, Ultrasound (MUI)
Date received	Sep 14, 2010
Decision date	Feb 16, 2011
Days to decision	155 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Dynarex Corporation</b>
Location	Orangeburg, NY, US
Contact	DANIEL CONSAGA
510(k) history	8 submissions · 8 cleared · 2008-2018

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k102637/> 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