

**K083467 REBOUND HRDV**Apr 23, 2009  
150 days to decisionK083467 · Product code: **FTL** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k083467/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Mesh, Surgical, Polymeric (FTL)
Date received	Nov 24, 2008
Decision date	Apr 23, 2009
Days to decision	150 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Minnesota Medical Development, Inc.</b>
Location	Stillwater, MN, US
Contact	STEVE NUSS
510(k) history	3 submissions · 3 cleared · 2007-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k083467/> 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 June 23, 2026