

**K080393 MODIFICATION TO REBOUND HRD**Mar 13, 2008  
29 days to decisionK080393 · Product code: **FTL** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k080393/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Mesh, Surgical, Polymeric (FTL)
Date received	Feb 13, 2008
Decision date	Mar 13, 2008
Days to decision	29 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	JULIE BULVER
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/k080393/> 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