

**K061892 PROPATCH SOFT TISSUE REPAIR MATRIX**Nov 22, 2006  
142 days to decisionK061892 · Product code: **FTM** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k061892/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Mesh, Surgical (FTM)
Date received	Jul 3, 2006
Decision date	Nov 22, 2006
Days to decision	142 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Cryolife, Inc.</b>
Location	Kennesaw, GA, US
Contact	JOHN D FERROS
510(k) history	12 submissions · 12 cleared · 2006-2019

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k061892/> 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 19, 2026