

**K961810 SUPPLE PERI-GUARD PERICARDIUM**Jun 13, 1996  
34 days to decisionK961810 · Product code: **FTM** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k961810/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mesh, Surgical (FTM)
Date received	May 10, 1996
Decision date	Jun 13, 1996
Days to decision	34 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Bio-Vascular, Inc.</b>
Location	St. Paul, MN, US
Contact	BRUCE A MACFARLANE
510(k) history	26 submissions · 25 cleared · 1986-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k961810/> 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 20, 2026