

**K063556 PERIPATCH ENDO-SLEEVE**Mar 20, 2007  
113 days to decisionK063556 · Product code: **FTM** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k063556/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mesh, Surgical (FTM)
Date received	Nov 27, 2006
Decision date	Mar 20, 2007
Days to decision	113 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Pm Devices, Inc.</b>
Location	Richmond, Bc, CA
Contact	TIM VERSPAGEN
510(k) history	4 submissions · 4 cleared · 2003-2007

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k063556/> 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 28, 2026