

**K081882 MODIFICATION TO XYLOS SURGICAL MESH**Jul 11, 2008  
9 days to decisionK081882 · Product code: **FTM** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k081882/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Mesh, Surgical (FTM)
Date received	Jul 2, 2008
Decision date	Jul 11, 2008
Days to decision	9 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Xylos Corporation</b>
Location	Washington, DC, US
Contact	GERRY ANN OSTER
510(k) history	9 submissions · 8 cleared · 1998-2011

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k081882/> 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 18, 2026