

**K071887 SPORTMESH OR ARTELON TISSUE  
REINFORCEMENT**Sep 18, 2007  
71 days to decision

K071887 · Product code: FTL · General &amp; Plastic Surgery

Source: <https://www.510kdatabase.net/k071887/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mesh, Surgical, Polymeric (FTL)
Date received	Jul 9, 2007
Decision date	Sep 18, 2007
Days to decision	71 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Artimplant AB</b>
Location	Washington, DC, US
Contact	TERRY S POWELL
510(k) history	7 submissions · 7 cleared · 2003-2007

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k071887/>. 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 19, 2026