

K061445 SURGIMESH WNFeb 1, 2007
253 days to decisionK061445 · Product code: **FTL** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k061445/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mesh, Surgical, Polymeric (FTL)
Date received	May 24, 2006
Decision date	Feb 1, 2007
Days to decision	253 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Aspide Medical
Location	Washington, DC, US
Contact	HOWARD M HOLSTEIN
510(k) history	4 submissions · 4 cleared · 2007-2012

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k061445/> Data sourced from FDA 510(k) public records (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. - Generated June 14, 2026