

K010228 GORE-TEX DUALMESH PLUS BIOMATERIAL WITH HOLESFeb 21, 2001
28 days to decisionK010228 · Product code: **FTL** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k010228/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Mesh, Surgical, Polymeric (FTL)
Date received	Jan 24, 2001
Decision date	Feb 21, 2001
Days to decision	28 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	W.L. Gore & Associates, Inc.
Location	McHenry, IL, US
Contact	R. LARRY PRATT
Website	http://www.gore.com/
510(k) history	163 submissions · 148 cleared · 1980-2025

W.L. Gore & Associates, Inc. is a global materials science company specializing in advanced medical devices. The company operates with a manufacturing facility in McHenry, US. The company has received FDA 510(k) clearances from total submissions since its first clearance in 1980. Cardiovascular devices represent a dominant category, including vascular grafts and balloon catheters. Recent clearances also span general surgery, plastic surgery, and gastroenterology applications. The latest FDA 510(k) clearance in 2025 reflects ongoing regulatory activity. W.L. Gore & Associa...

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Device record: <https://www.510kdatabase.net/k010228/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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