

K061030 BIOBLANKET SURGICAL MESHMay 9, 2006
25 days to decisionK061030 · Product code: **FTM** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k061030/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Mesh, Surgical (FTM)
Date received	Apr 14, 2006
Decision date	May 9, 2006
Days to decision	25 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Kensey Nash Corporation
Location	Exton, PA, US
Contact	DEBORAH RACIOPPI
510(k) history	6 submissions · 6 cleared · 2006-2013

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k061030/> 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 May 19, 2026