

K936186 AESCULAP SKIN GRAFT MESHERMar 10, 1994
77 days to decisionK936186 · Product code: **FZW** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k936186/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Expander, Surgical, Skin Graft (FZW)
Date received	Dec 23, 1993
Decision date	Mar 10, 1994
Days to decision	77 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Aesculap, Inc.
Location	Burlingame, CA, US
Contact	MARY ELLEN HOLDEN
510(k) history	207 submissions · 201 cleared · 1991-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k936186/> 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 28, 2026