FDA 510(k) Premarket Notification

CLEARED

K936186 AESCULAP SKIN GRAFT MESHER

Mar 10, 1994
77 days to decision

K936186 · Product code: **FZW** · General & Plastic Surgery
Source: 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