

K922576 AESCULAP MANUAL ENDOSCOPIC INSTRUMENTSJun 10, 1993
374 days to decisionK922576 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k922576/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 1, 1992
Decision date	Jun 10, 1993
Days to decision	374 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k922576/> 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