

**K014207 REPOSABLE INSTRUMENT SYSTEM, MODELS
MF001-MF083**Mar 21, 2002
90 days to decisionK014207 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k014207/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Dec 21, 2001
Decision date	Mar 21, 2002
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

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