

K951353 HYBRID SURGICAL DEVICES, MODIFICIATIONApr 6, 1995
38 days to decisionK951353 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k951353/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Feb 27, 1995
Decision date	Apr 6, 1995
Days to decision	38 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Laser Centers of America, Inc.
Location	Cincinnati, OH, US
Contact	RICHARD L STUDER
510(k) history	1 submissions · 1 cleared · 1995-1995

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k951353/> 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 July 6, 2026