

K924410 LAPAROSCOPIC/ENDOSCOPIC INSTRUMENTATIONJun 25, 1993
298 days to decisionK924410 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k924410/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Aug 31, 1992
Decision date	Jun 25, 1993
Days to decision	298 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Core Dynamics, Inc.
Location	Jacksonville, FL, US
Contact	BILL DENNIS
510(k) history	23 submissions · 23 cleared · 1990-1999

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k924410/> 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 29, 2026