FDA 510(k) Premarket Notification

K924410 LAPAROSCOPIC/ENDOSCOPIC INSTRUMENTATION

CLEARED

Jun 25, 1993
298 days to decision

K924410 · Product code: **GEI** · General & Plastic Surgery
Source: 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