

K960009 STORZ S2050 SERIES BIPOLAR FORCEPSMar 6, 1996
64 days to decisionK960009 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k960009/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jan 2, 1996
Decision date	Mar 6, 1996
Days to decision	64 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Storz Instrument Co.
Location	Mchenry, IL, US
Contact	AUDREY SWEARINGEN
510(k) history	101 submissions · 100 cleared · 1976-1998

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k960009/> 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 May 21, 2026