FDA 510(k) Premarket Notification

K960009 STORZ S2050 SERIES BIPOLAR FORCEPS

CLEARED

Mar 6, 1996
64 days to decision

K960009 · Product code: **GEI** · General & Plastic Surgery
Source: 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