

K091440 LAPAROSCOPIC RESECTION DEVICE, MODEL SR326LOct 9, 2009
147 days to decisionK091440 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k091440/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	May 15, 2009
Decision date	Oct 9, 2009
Days to decision	147 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Bovie Medical
Location	St. Petersburg, FL, US
Contact	RICHARD KOZLOFF
510(k) history	8 submissions · 8 cleared · 2006-2010

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k091440/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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