

**K082568 SALINE ENHANCED ELECTROSURGICAL RESECTION
(SEER) DEVICE**Nov 17, 2008
74 days to decisionK082568 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k082568/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Sep 4, 2008
Decision date	Nov 17, 2008
Days to decision	74 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k082568/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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