

**K920217 MAX-E-VAC**Jul 23, 1992  
189 days to decisionK920217 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k920217/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jan 16, 1992
Decision date	Jul 23, 1992
Days to decision	189 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Leisegang Medical, Inc.</b>
Location	Miami, FL, US
Contact	DOUGLAS KWART
510(k) history	10 submissions · 10 cleared · 1986-1999

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k920217/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 29, 2026