

**K000936 VAPR 2.3MM WEDGE ELECTRODE FOR USE WITH VAPR SYSTEM**

Apr 19, 2000  
27 days to decision

K000936 · Product code: **GEI** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k000936/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Mar 23, 2000
Decision date	Apr 19, 2000
Days to decision	27 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Mitek Products</b>
Location	Westwood, MA, US
Contact	MARY P LEGRAW
510(k) history	30 submissions · 30 cleared · 1996-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k000936/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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