

**K012695 VANGUARD REPROCESSED ARTHROSCOPIC  
WANDS**Oct 30, 2001  
77 days to decisionK012695 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k012695/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Aug 14, 2001
Decision date	Oct 30, 2001
Days to decision	77 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Vanguard Medical Concepts, Inc.</b>
Location	Plant City, FL, US
Contact	MIKE SAMMON
510(k) history	33 submissions · 33 cleared · 1991-2005

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k012695/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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