

**K201164 Venus Viva MD Device**Jun 26, 2020  
56 days to decisionK201164 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k201164/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	May 1, 2020
Decision date	Jun 26, 2020
Days to decision	56 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Venus Concept USA, Inc.</b>
Location	Sunrise, FL, US
Contact	Yoni Iger
510(k) history	11 submissions · 11 cleared · 2014-2022

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k201164/> 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 May 26, 2026