

**K131696 VESALIUS**Jan 31, 2014  
235 days to decisionK131696 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k131696/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 10, 2013
Decision date	Jan 31, 2014
Days to decision	235 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Telea Electronic Engineering Srl</b>
Location	Imola (Bo), IT
Contact	TESTA MARISA
510(k) history	2 submissions · 2 cleared · 2014-2016

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k131696/> 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 3, 2026