

**K952973 THE TULIP MONO-POLAR CAUTERY SPATULA**Sep 11, 1995  
76 days to decisionK952973 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k952973/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 27, 1995
Decision date	Sep 11, 1995
Days to decision	76 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>The Tulip Mfg. Co.</b>
Location	Mesa, AZ, US
Contact	RICHARD S HUNTER
510(k) history	24 submissions · 24 cleared · 1995-1996

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k952973/> 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 July 1, 2026