

**K955501 THE TULIP MONO POLAR ROTATABLE ALLIGATOR  
FORCEP**Jan 17, 1996  
47 days to decisionK955501 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k955501/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Dec 1, 1995
Decision date	Jan 17, 1996
Days to decision	47 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k955501/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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