

**K014260 TISSUELINK SOLID CYLINDER MONOPOLAR DEVICE**

Mar 27, 2002  
90 days to decision

K014260 · Product code: **GEI** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k014260/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Dec 27, 2001
Decision date	Mar 27, 2002
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Tissuelink Medical, Inc.</b>
Location	Dover, NH, US
Contact	VICKI ANASTASI
510(k) history	12 submissions · 12 cleared · 2000-2008

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k014260/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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