

**K010192 ENDOSCOPIC TUBE SHAFT INSTRUMENTS FOR  
MONOPLAR COAGULATION**Apr 2, 2001  
70 days to decisionK010192 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k010192/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jan 22, 2001
Decision date	Apr 2, 2001
Days to decision	70 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Antyllos Medizintechnik GmbH</b>
Location	Wurmlingen, DE
Contact	HARALD JUNG
510(k) history	1 submissions · 1 cleared · 2001-2001

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k010192/> 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 3, 2026