

**K982472 GENICON TROCAR**Feb 4, 1999  
204 days to decisionK982472 · Product code: **GEA** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k982472/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Surgical, General & Plastic Surgery (GEA)
Date received	Jul 15, 1998
Decision date	Feb 4, 1999
Days to decision	204 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Genicon, LC</b>
Location	Orlando, FL, US
Contact	GARY W HABERLAND
510(k) history	6 submissions · 6 cleared · 1999-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k982472/> 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 29, 2026