

**K161307 NICO Myriad**Dec 21, 2016  
225 days to decisionK161307 · Product code: **GEI** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k161307/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	May 10, 2016
Decision date	Dec 21, 2016
Days to decision	225 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Nico Corporation</b>
Location	Indianapolis, IN, US
Contact	SEAN SPENCE
510(k) history	9 submissions · 9 cleared · 2012-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k161307/> 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 May 27, 2026