

K182340 NICO MyriadSep 20, 2018
23 days to decisionK182340 · Product code: **GEI** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k182340/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Aug 28, 2018
Decision date	Sep 20, 2018
Days to decision	23 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Nico Corporation
Location	Indianapolis, IN, US
Contact	Sean Spence
510(k) history	9 submissions · 9 cleared · 2012-2024

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k182340/> Data sourced from FDA 510(k) public records (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. - Generated May 27, 2026