FDA 510(k) Premarket Notification

# K161307 NICO Myriad

**CLEARED**

Dec 21, 2016
225 days to decision

K161307 · Product code: **GEI** · Obstetrics & Gynecology
Source: 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

| | |
|---|---|
| Company | **Nico Corporation** |
| Location | Indianapolis, IN, US |
| Contact | SEAN SPENCE |
| 510(k) history | 9 submissions · 9 cleared · 2012-2024 |