FDA 510(k) Premarket Notification

K182340 NICO Myriad

CLEARED

Sep 20, 2018
23 days to decision

K182340 · Product code: **GEI** · Obstetrics & Gynecology
Source: 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