FDA 510(k) Premarket Notification

K120151 NEBULAE I

CLEARED

Aug 17, 2012
212 days to decision

K120151 · Product code: **HIF** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k120151/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Insufflator, Laparoscopic (HIF)
Date received	Jan 18, 2012
Decision date	Aug 17, 2012
Days to decision	212 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Northgate Technologies, Inc.**
Location	Arlington Heights, IL, US
Contact	CASEY KUREK
510(k) history	55 submissions · 55 cleared · 1991-2021