

K120151 NEBULAE IAug 17, 2012
212 days to decisionK120151 · Product code: **HIF** · Obstetrics & GynecologySource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k120151/> 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 14, 2026