

**K914401 UNIVERSAL GENERAL USE KIT #7-510-16**Nov 27, 1991  
56 days to decisionK914401 · Product code: **HIF** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k914401/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Insufflator, Laparoscopic (HIF)
Date received	Oct 2, 1991
Decision date	Nov 27, 1991
Days to decision	56 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Northgate Technologies, Inc.</b>
Location	Arlington Heights, IL, US
Contact	PETER A MANZIE
510(k) history	55 submissions · 55 cleared · 1991-2021

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k914401/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 28, 2026