FDA 510(k) Premarket Notification

K914402 RECIRCULATION KIT #7-510-18

CLEARED

Nov 27, 1991
56 days to decision

K914402 · Product code: **HIF** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k914402/

SUBMISSION DETAILS

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

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k914402/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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