

**K060751 INOVO, INC. ACCUPULSE SINGLE LUMEN
CONSERVING REGULATOR**Jul 18, 2006
119 days to decisionK060751 · Product code: **NFB** · Anesthesiology
Source: <https://www.510kdatabase.net/k060751/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Conserver, Oxygen (NFB)
Date received	Mar 21, 2006
Decision date	Jul 18, 2006
Days to decision	119 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Inovo, Inc.
Location	Great Neck, NY, US
Contact	MICHAEL T DILDINE
510(k) history	6 submissions · 6 cleared · 2003-2014

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k060751/> 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 24, 2026