

**K131023 THERAFLOW SINGLE LUMEN AND THERAFLOW
DUAL LUMEN**Oct 11, 2013
182 days to decisionK131023 · Product code: **NFB** · Anesthesiology
Source: <https://www.510kdatabase.net/k131023/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Conservor, Oxygen (NFB)
Date received	Apr 12, 2013
Decision date	Oct 11, 2013
Days to decision	182 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Medical Products, LLC
Location	Indianapolis, IN, US
Contact	DAVID MCGILL
510(k) history	1 submissions · 1 cleared · 2013-2013

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k131023/> 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