FDA 510(k) Premarket Notification

K060751 INOVO, INC. ACCUPULSE SINGLE LUMEN CONSERVING REGULATOR

CLEARED

Jul 18, 2006
119 days to decision

K060751 · 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