FDA 510(k) Premarket Notification

K882328 HI FLOW 50

CLEARED

Jun 28, 1988
22 days to decision

K882328 · Product code: **BYJ** · Anesthesiology
Source: https://www.510kdatabase.net/k882328/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Liquid-oxygen, Portable (BYJ)
Date received	Jun 6, 1988
Decision date	Jun 28, 1988
Days to decision	22 days
Third-party review	No

APPLICANT

Company	**Minnesota Valley Engineering, Inc.**
Location	New Prague, MN, US
Contact	SCHOENBAUER
510(k) history	4 submissions · 4 cleared · 1987-1988