FDA 510(k) Premarket Notification

K875352 HI FLOW PORTABLE LIQUID OXYGEN UNIT

CLEARED

Feb 29, 1988
60 days to decision

K875352 · Product code: **BYJ** · Anesthesiology
Source: https://www.510kdatabase.net/k875352/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Liquid-oxygen, Portable (BYJ)
Date received	Dec 31, 1987
Decision date	Feb 29, 1988
Days to decision	60 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