FDA 510(k) Premarket Notification

K863978 PORTABLE LIQUID OXYGEN UNIT (MODIFICATION)

CLEARED

Dec 15, 1986
62 days to decision

K863978 · Product code: **BYJ** · Anesthesiology
Source: https://www.510kdatabase.net/k863978/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Liquid-oxygen, Portable (BYJ)
Date received	Oct 14, 1986
Decision date	Dec 15, 1986
Days to decision	62 days
Third-party review	No

APPLICANT

Company	**Penox Technologies, Inc.**
Location	Pittston, PA, US
Contact	THANA A FRANCE
510(k) history	10 submissions · 10 cleared · 1985-1987