FDA 510(k) Premarket Notification

K013931 OMNI OXYGEN SYSTEM, MODEL 1000

CLEARED

Mar 27, 2002
119 days to decision

K013931 · Product code: **CAW** · Anesthesiology
Source: https://www.510kdatabase.net/k013931/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Generator, Oxygen, Portable (CAW)
Date received	Nov 28, 2001
Decision date	Mar 27, 2002
Days to decision	119 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Sequal Technologies, Inc.**
Location	San Diego, CA, US
Contact	PAM JACKSON
510(k) history	4 submissions · 4 cleared · 2001-2009