FDA 510(k) Premarket Notification

K131990 OXYGEN GENERATOR LIQUEFIER - (OGL)

CLEARED

Feb 21, 2014
238 days to decision

K131990 · Product code: **CAW** · Anesthesiology
Source: https://www.510kdatabase.net/k131990/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Generator, Oxygen, Portable (CAW)
Date received	Jun 28, 2013
Decision date	Feb 21, 2014
Days to decision	238 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Essex Cryogenics of Missouri, Inc.**
Location	St. Louis, MO, US
Contact	PAUL DRYDEN
510(k) history	5 submissions · 5 cleared · 1988-2014