FDA 510(k) Premarket Notification

K001121 OXYGEN APPLIED SYMPTOM IMPROVEMENT SYSTEM (OASIS)

CLEARED

Nov 29, 2001
601 days to decision

K001121 · Product code: **CBF** · Anesthesiology
Source: https://www.510kdatabase.net/k001121/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Chamber, Hyperbaric (CBF)
Date received	Apr 7, 2000
Decision date	Nov 29, 2001
Days to decision	601 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Engineered Medical Technology, Inc.**
Location	Victoria, TX, US
Contact	DON SCHMIELEY
510(k) history	1 submissions · 1 cleared · 2001-2001