

**K001121 OXYGEN APPLIED SYMPTOM IMPROVEMENT  
SYSTEM (OASIS)**Nov 29, 2001  
601 days to decisionK001121 · Product code: **CBF** · Anesthesiology  
Source: <https://www.510kdatabase.net/k001121/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k001121/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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