

**K813436 OXYCON**Jan 7, 1982  
30 days to decisionK813436 · Product code: **CAW** · Anesthesiology  
Source: <https://www.510kdatabase.net/k813436/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Generator, Oxygen, Portable (CAW)
Date received	Dec 8, 1981
Decision date	Jan 7, 1982
Days to decision	30 days
Third-party review	No

**APPLICANT**

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Company	<b>Home Medical Mfg., Co., Inc.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1982-1982

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Device record: <https://www.510kdatabase.net/k813436/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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