

**K885079 HORIZON 2000 TRANSCUTANEOUS GAS #26X-  
OPT-275**Feb 17, 1989  
71 days to decisionK885079 · Product code: **LPP** · Anesthesiology  
Source: <https://www.510kdatabase.net/k885079/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Oxygen, Cutaneous, For Uses Other Than For Infant Not Under Gas Anesthesia (LPP)
Date received	Dec 8, 1988
Decision date	Feb 17, 1989
Days to decision	71 days
Third-party review	No

**APPLICANT**

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Company	<b>Mennen Medical, Inc.</b>
Location	Clarence, NY, US
Contact	RONALD A WIDMAN
510(k) history	34 submissions · 34 cleared · 1985-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k885079/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 21, 2026