

**K895274 HYPEX**Feb 23, 1990  
184 days to decisionK895274 · Product code: **BYW** · AnesthesiologySource: <https://www.510kdatabase.net/k895274/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Rebreathing (BYW)
Date received	Aug 23, 1989
Decision date	Feb 23, 1990
Days to decision	184 days
Third-party review	No

**APPLICANT**

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Company	<b>Global Business, Inc.</b>
Location	Midlothian, VA, US
Contact	LENKEY III
510(k) history	1 submissions · 1 cleared · 1990-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k895274/>; 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 26, 2026