

**K952084 HUMID-VENT FILTER PEDI**Jun 12, 1995  
40 days to decisionK952084 · Product code: **BYD** · Anesthesiology  
Source: <https://www.510kdatabase.net/k952084/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Condenser, Heat And Moisture (artificial Nose) (BYD)
Date received	May 3, 1995
Decision date	Jun 12, 1995
Days to decision	40 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Gibeck, Inc.</b>
Location	Indianapolis, IN, US
Contact	BRIAN GRIGSBY
510(k) history	15 submissions · 15 cleared · 1990-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k952084/> 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