

**K925622 FHB, FHB-T, FHBFLEX, FHBFLEX-T**May 26, 1993  
201 days to decisionK925622 · Product code: **BYD** · Anesthesiology  
Source: <https://www.510kdatabase.net/k925622/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Condenser, Heat And Moisture (artificial Nose) (BYD)
Date received	Nov 6, 1992
Decision date	May 26, 1993
Days to decision	201 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Icor AB</b>
Location	Bromma, SE
Contact	ANDRAS GEDEON
510(k) history	16 submissions · 16 cleared · 1986-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k925622/> 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