

**K951652 VENTIMASK MK IV**Apr 27, 1995  
17 days to decisionK951652 · Product code: **BYF** · Anesthesiology  
Source: <https://www.510kdatabase.net/k951652/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mask, Oxygen, Low Concentration, Venturi (BYF)
Date received	Apr 10, 1995
Decision date	Apr 27, 1995
Days to decision	17 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Flexicare Medical, Ltd.</b>
Location	Somerset, GB
Contact	COLIN BECKETT
510(k) history	7 submissions · 7 cleared · 1995-2015

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k951652/> 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 24, 2026