

**K960587 CPROTECTOR 2000**Jul 21, 1997  
525 days to decisionK960587 · Product code: **CBP** · Anesthesiology  
Source: <https://www.510kdatabase.net/k960587/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Valve, Non-rebreathing (CBP)
Date received	Feb 12, 1996
Decision date	Jul 21, 1997
Days to decision	525 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Certified Safety Mfg., Inc.</b>
Location	Kansas City, MO, US
Contact	HOWARD GERSON
510(k) history	17 submissions · 14 cleared · 1994-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k960587/> 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 25, 2026