

**K810348 CM 3**Feb 23, 1981  
13 days to decisionK810348 · Product code: **CBP** · AnesthesiologySource: <https://www.510kdatabase.net/k810348/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Valve, Non-rebreathing (CBP)
Date received	Feb 10, 1981
Decision date	Feb 23, 1981
Days to decision	13 days
Third-party review	No

**APPLICANT**

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Company	<b>Instrumentation Industries, Inc.</b>
Location	Mchenry, IL, US
510(k) history	45 submissions · 44 cleared · 1977-2019

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k810348/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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