

**K791850 ONE-WAY VALVE-FEMALE, M-503**Oct 1, 1979  
12 days to decisionK791850 · Product code: **CBP** · Anesthesiology  
Source: <https://www.510kdatabase.net/k791850/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Valve, Non-rebreathing (CBP)
Date received	Sep 19, 1979
Decision date	Oct 1, 1979
Days to decision	12 days
Third-party review	No

**APPLICANT**

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Company	<b>Mem Medical, Inc.</b>
Location	Mchenry, IL, US
510(k) history	27 submissions · 27 cleared · 1979-1979

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

Device record: <https://www.510kdatabase.net/k791850/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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