

**K896500 CWD/PPG MODULE**Jul 18, 1990  
246 days to decisionK896500 · Product code: **JOM** · Cardiovascular  
Source: <https://www.510kdatabase.net/k896500/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plethysmograph, Photoelectric, Pneumatic Or Hydraulic (JOM)
Date received	Nov 14, 1989
Decision date	Jul 18, 1990
Days to decision	246 days
Third-party review	No

**APPLICANT**

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Company	<b>Life Sciences Manufacturing, Inc.</b>
Location	Mchenry, IL, US
Contact	ROBERT S CUTLER
510(k) history	12 submissions · 12 cleared · 1979-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k896500/>, 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 June 29, 2026