



# Current Good Manufacturing Practices: Pharmaceutical, Biologics, and Medical Device Regulations and Guidance Documents Concise Reference

By Mindy J. Allport-Settle

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### **Editorial Review**

#### **About the Author**

Mindy Allport-Settle has served as a key executive, board member, and consultant for some of the best companies in the pharmaceutical and FDA-regulated industry. Since joining PharmaLogika as CEO in 2008 and over her career, she has provided informed guidance in regulatory compliance, corporate structuring, restructuring and turnarounds, new drug submissions, research & development and product commercialization strategies, operational, project and contract management, and new business development. Her experience and dedication have resulted in international recognition as the developer of the only FDA-recognized and benchmarked quality systems training and development business methodology designed for regulated industries. Her education includes a Bachelor's degree from the University of North Carolina, an MBA in Global Management from the University of Phoenix, and completion of the corporate governance course series in audit committees, compensation committees, and board effectiveness at Harvard Business School.

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