

# Dietary Supplements: Current Good Manufacturing Practice, Labeling and Premarket Notification Concise Reference

Mindy J. Allport-Settle

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This book is designed to be a unified reference source for the U.S. Food and Drug Administration's Dietary Supplements regulations, guidance, and associated documents. Dietary Supplements Regulations and Guidance: \* FDA Overview and Orientation \* Introduction to Dietary Supplements \* Part I: Regulations -Part 111: Current Good Manufacturing Practice in Manufacturing, Packaging, Labeling, or Holding Operations for Dietary Supplements - Part 101: Food Labeling - Part 190: Dietary Supplements \* Part II: Legislative Acts and Significant Amendments - Dietary Supplement and Nonprescription Drug Consumer Protection Act - Dietary Supplement Health and Education Act of 1994 \* Part III: Guidance and Associated Documents - Dietary Supplements Final Rule - Dietary Supplement Current Good Manufacturing Practices (CGMPs) and Interim Final Rule (IFR) Facts - Dietary Supplement Labeling Guide - Structure/Function Claims, Small Entity Compliance Guide - Notification of a Health Claim or Nutrient Content Claim Based on an Authoritative Statement of a Scientific Body - Questions and Answers Regarding Adverse Event Reporting and Recordkeeping for Dietary Supplements as Required by the Dietary Supplement and Nonprescription Drug Consumer Protection Act - Responses to Questions about Codex and Dietary Supplements Reference Tools \* Part IV: Combined Glossary and Index

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