



GMP TRAINING



Good Manufacturing Practices Overview for Management: NHPs Health Canada NHPD Good Manufacturing Practices (GMPs)

2 HOURS
Management
Overview



WHO SHOULD ATTEND

- Executives in the natural health product (NHP) industry
- Non-technical audience
- Managers with no formal training in GMP

PURPOSE OF THE WORKSHOP

- Provides a fast-paced learning package that enables you to gain a working knowledge of GMP
- Shows you how to apply GMP principles
- Simplifies the regulations and helps you understand why you need to comply and what you need to do to comply
- Identifies Health Canada NHPD's regulatory expectations
- Shows you how to implement GMP to avoid surprises in government audits
- Greatly reduces the risk of failing an audit
- Helps you prepare an action plan for implementing GMP immediately on your return to the workplace
- Employs interactive Mr. GMP Exercise™
- Answers your questions based on the most current interpretations of the GMP regulations
- Gives you a participant binder, marked quiz, course certificate and indexed GMP pocket booklet. The Green Book™ will be an indispensable reference for your day-to-day activities in the workplace.

WORKSHOP OBJECTIVES

- Acquire and apply the basic concepts of GMP
- Recognize the importance of GMP compliance
- Understand the role of Quality Control

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GMP Overview for Management: NHPs ...continued

WHAT YOU WILL LEARN

1. What is GMP?
 - a. Brief history of GMP
 - b. Why is GMP important?
 - c. What is a NPN?
2. Role of Quality (QA/QC)
 - a. Quality Assurance vs. Quality Control
 - b. What does Quality Ownership mean?
 - c. What is SISPO?
3. Equipment
 - a. Cleaning requirements
 - b. Calibrate what and when?
 - c. PM programs
4. Validation
 - a. What is Validation?
 - b. What types of validation are required?
 - c. How do you perform validation?
5. Sanitation and Hygiene
 - a. Sanitation Programs
 - b. How do you prevent contamination?
 - c. Personal hygiene
6. Labeling
 - a. What needs to be on a label?
 - b. Identifying product status
 - c. What does 'Quarantined' mean?
7. Records and Documentation
 - a. What is a Record?
 - b. Good Documentation Practices (GDP)
 - c. Common documentation errors
8. Procedures
 - a. SOPs
 - b. Regulatory requirements
 - c. How do you control these?
9. Deviations/Change Control
 - a. Why is it important to document Deviations?
 - b. 1/10/100 Rule
 - c. What is Change Control?
10. The Manufacturing Cycle
 - a. Pre-Manufacturing Activities
 - b. What are some specific GMP controls?
 - c. Consequences of poor controls
11. Audits
 - a. Risk classifications
 - b. How to prepare for a successful audit
 - c. What do Health Canada inspectors look for?