

MEET YOUR GMP TRAINING NEEDS



COURSE	Fundamentals of Good Manufacturing Practices: Drugs	GMP Overview for Management: Drugs	Drug GMP for Importers and Distributors
DURATION	1 day	2 hours	½ day
WHO SHOULD ATTEND	Fabricators Packagers/Labelers Testers	Managers and executives Non-technical	Importers Distributors
REGULATIONS	Health Canada HPFBI GMP and/or FDA Title 21, Food & Drugs Parts 11, 210 and 211	Health Canada HPFBI GMP and/or FDA Title 21, Food & Drugs Parts 11, 210 and 211	Health Canada HPFBI GMP
	Combination Drug and Natural Health Product/Dietary Supplement Courses Available		
DESCRIPTION	<ul style="list-style-type: none"> • Gain a basic understanding of GMP/cGMP regulations • Review a brief history of GMP; understand the role of quality control and quality assurance; learn about equipment use, validation, sanitation, labeling, records, records and good documentation; and look at deviations, change control and audits • Through the experiential Mr. GMP Exercise™, learn how to apply GMP/cGMP regulations • Implementations are geared to your audience, such as employees new to GMP/cGMP, or experienced employees needing to stay current with GMP/cGMP. On completion, learners meet HPFBI or FDA training requirements (as per HPFBI GMP regulations C.02.006) • We train your staff on-site (closed sessions for 6 to 30 participants). Audiovisual equipment available upon request 		
COMMON LEARNING OBJECTIVES	<ul style="list-style-type: none"> • Understand and apply the basic concepts of GMP/cGMP regulations, recognizing the importance of GMP/cGMP compliance • Follow GMP/cGMP as they relate to the specific audience (fabricators, packagers/labelers, testers, importers, distributors and wholesalers) • Understand the role of Quality Control, particularly in testing and release activities • Follow Good Documentation Practices • Apply the regulations to prepare for a successful audit 		



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