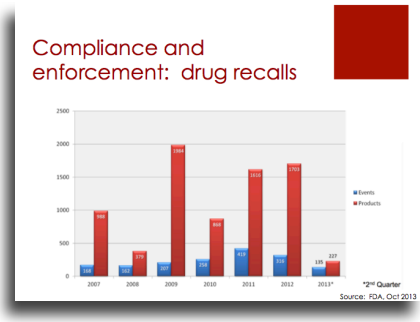
GMP Update

An in-house course from *LearningPlus*

Keep your quality, technical, and management groups current with what is happening in regards to pharmaceutical GMPs, compliance, and quality.



Description

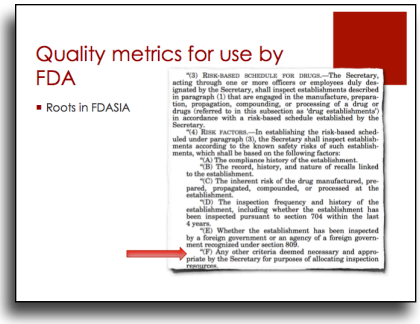
Each year, LearningPlus revises its popular ***GMP Update*** course with information about new and revised regulations, guidances, and expectations from the FDA and regulatory agencies in Canada and Europe.

Designed for quality staff, support professionals, and management, this 3-hour workshop also includes compliance and inspection trends, cutting-edge topics, and a case study where participants can apply their knowledge and critical "GMP thinking" skills.

Recent topics have included data integrity, quality metrics and measurement, Warning Letter reviews, Ebola clinical trials, and "track and trace".

LearningPlus provides on-site GMP training that can be adapted to your specific needs.

Contact Jim Vesper at +1 585.442.0170 or email jvesper@learningplus.com.



Course goal

Provide a summary of important changes in pharmaceutical regulations, expectations, and trends in compliance, inspectional, and quality issues.

Audience

Those in management, quality, development, compliance, laboratory, operations, and other technical areas.

Objectives

- Discuss changes in GMP-related regulations, requirements, and guidances
- Describe trends in related compliance activities that have been published by regulatory agencies
- Examine particular Warning Letters and compliance activities of interest
- Discuss examples and cases that emphasize the importance of quality systems and GMP thinking

Length

Approximately 3 hours. (Two sessions can be given in one day.)