

# Assignment

1. Explain the importance of Current Good Manufacturing Practices (cGMP) and discuss how it impacts pharmaceutical manufacturing.
2. Describe the key elements of cGMP, focusing on their role in maintaining product quality and patient safety.
3. What are the regulatory frameworks governing cGMP in different regions? Compare the roles of USFDA, WHO, and EMA in ensuring compliance.
4. Discuss the role of documentation in cGMP. How do Master Production Records (MPR) and Batch Records contribute to compliance?
5. Highlight the modern trends in cGMP, such as continuous manufacturing, and explain its advantages in the pharmaceutical industry.