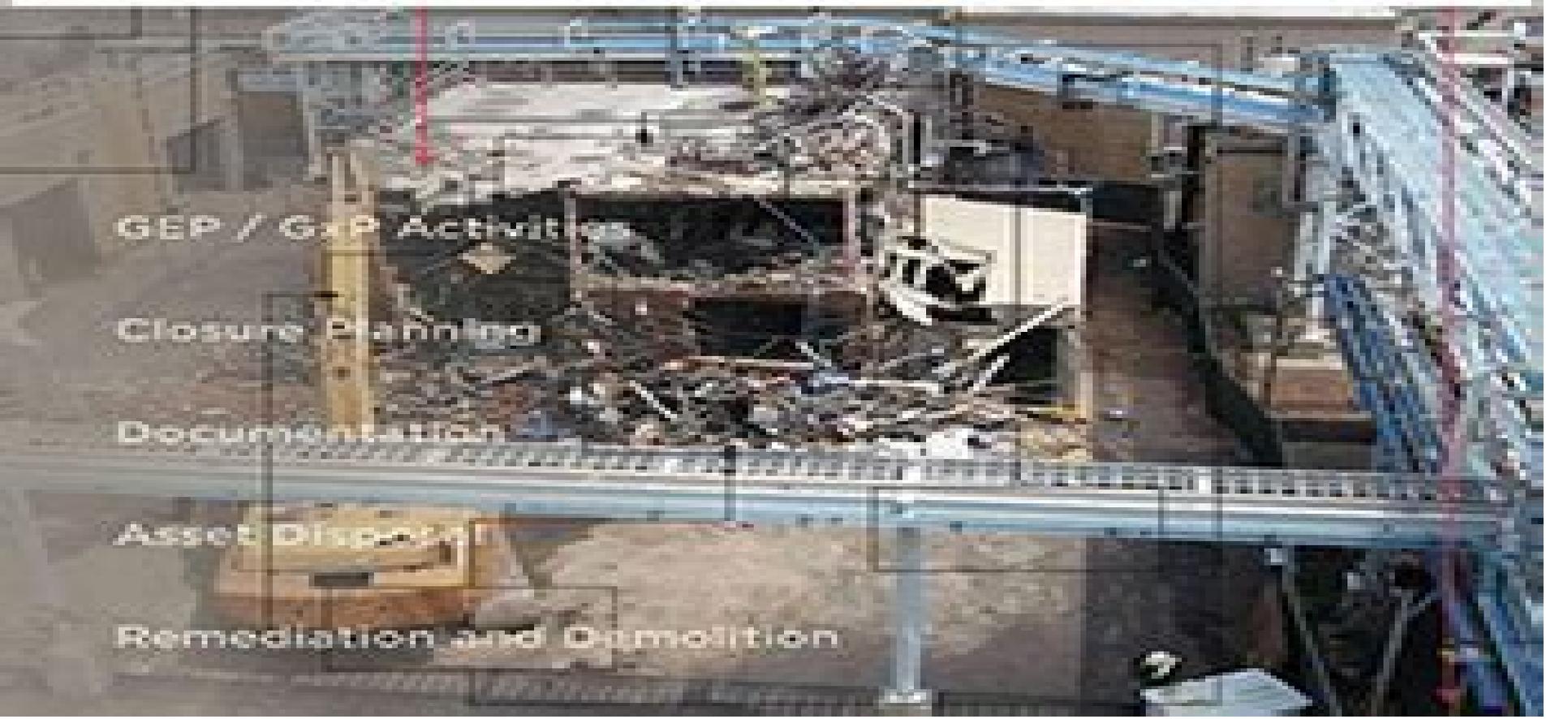


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The background image shows an industrial facility, likely a pharmaceutical plant, in the process of being demolished. Large pieces of machinery and structural elements are visible, with a significant amount of debris on the ground. The scene is captured from an elevated perspective, showing the complex network of pipes and structural beams.

GEP / GxP Activities

Closure Planning

Documentation

Asset Disposal

Remediation and Demolition

# Good Practice Decommissioning Pharma Ispe

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## **Good Practice Decommissioning Pharma Ispe:**

*ISPE Good Practice Guide Ispe*,2017-06-27      **Equipment Qualification in the Pharmaceutical Industry** Steven Ostrove,2019-06-13 Equipment Qualification in the Pharmaceutical Industry provides guidance and basic information for the preparation of a quality qualification program It has been noted that there is a general lack of understanding in the industry especially for those new to the industry as to what constitutes a compliant qualification program Even experienced professionals have felt a lack of security in reaching a compliant state This book outlines a guideline for the preparation and execution of qualification protocols including the installation IQ operational OQ and performance PQ protocols It discusses the importance of related qualification programs e g quality systems commissioning computer system and cleaning and how to incorporate them into a fully compliant qualification program Furthermore it provides matrices of what could be included in each type of protocol for major types of process equipment While primarily for people entering the pharmaceutical industry those established in the field will benefit from the multiple examples and matrices as well as integration of related systems Equipment Qualification in the Pharmaceutical Industry provides students and pharmaceutical scientists a guideline for the preparation and execution of qualification installation operational and performance protocols Incorporates good manufacturing processes into a compliant qualification program Provides examples of protocol layout Includes matrices for major process equipment installation quality operational quality and performance quality requirements      Pharmaceutical Manufacturing Handbook Shayne Cox Gad,2008-03-11 With its coverage of Food and Drug Administration regulations international regulations good manufacturing practices and process analytical technology this handbook offers complete coverage of the regulations and quality control issues that govern pharmaceutical manufacturing In addition the book discusses quality assurance and validation drug stability and contamination control all key aspects of pharmaceutical manufacturing that are heavily influenced by regulatory guidelines The team of expert authors offer you advice based on their own firsthand experience in all phases of pharmaceutical manufacturing      *Pharmaceutical Computer Systems Validation* Guy Wingate,2016-04-19 Thoroughly revised to include the latest industry developments the Second Edition presents a comprehensive overview of computer validation and verification principles and how to put them into practice To provide the current best practice and guidance on identifying and implementing improvements for computer systems the text extensively reviews regulations of pharmaceuticals healthcare products blood processing medical devices clinical systems and biotechnology Ensuring that organizations transition smoothly to the new system this guide explains how to implement the new GMP paradigm while maintaining continuity with current practices In addition all 24 case studies from the previous edition have been revised to reflect the new system      **Ensuring the Integrity of Electronic Health Records** Orlando López,2020-12-21 Data integrity is a critical aspect to the design implementation and usage of any system which stores processes or retrieves data The overall intent of any data integrity technique is the same ensure data is recorded exactly as

intended and upon later retrieval ensure the data is the same as it was when originally recorded Any alternation to the data is then traced to the person who made the modification The integrity of data in a patient s electronic health record is critical to ensuring the safety of the patient This book is relevant to production systems and quality control systems associated with the manufacture of pharmaceuticals and medical device products and updates the practical information to enable better understanding of the controls applicable to e records The book highlights the e records suitability implementation and associated risk assessed controls and e records handling The book also provides updated regulatory standards from global regulatory organizations such as MHRA Medicines and Healthcare Products Regulatory Agency UK FDA Food and Drug Administration US National Medical Products Association China TGA Therapeutic Goods Administration Australia SIMGP Russia State Institute of Medicines and Good Practices and the World Health Organization to name a few

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