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1. The most important Utilities in the pharmaceutical Manufacturing site, what should the layout and design of Pharmaceutical Manufacturing Site aim to and illustrate simple layout for material and people flow.
2. Dosage form: what are the needs for dosage forms, classify the dosage forms according to "physical form and route of administration" and illustrate simple diagram for one of dosage forms.
3. Process validation, what is the purpose of process validation, what should be validated and differentiate between New FPPs and Established FPPs in Process validation & evaluation.
4. ISO 19001 and ISO 14001; overview, benefits, comparison and what are the differences between certification and accreditation.
5. According to GMP guidelines please give the definition for the three types of deficiency, manufacturing process Information required, what are the steps and please give us the needs of scale Up Data.