

GMP Workshop on construction and renovation of pharmaceutical manufacturing sites

- Understand the criticality of construction and renovation work to GMP status of pharmaceutical premises
- Inform yourself of regulatory requirements for premises and equipment
- Gain fundamentals of the management of construction or renovation from a manufacturer's perspective
- Gain fundamentals of the management of construction or renovation from the authority's viewpoint
- Define design and layout for manufacturing on non-sterile products
- Gain fundamentals of the design and specification of facilities (e.g. material flow, personnel flow, surfaces)
- Understand the impact of construction and renovations on HVAC systems
- Understand the criticality of procurement, installation and qualification of GMP relevant equipment

Join us for pratical sessions, theoretical background and professional exchange!

Venue: Conference hall + practical sessions on site

Trainers: experienced trainers from international pharma industry

Duration: 4-5 days

Target group: professions from pharmaceutical manufacturers (engineers, heads of Quality Assurance or heads of production), NMRA



E-Mail: irmgard.buchkremer@medeor.de