February **25–26**, 2027 | **Amsterdam**, Netherlands | **#EAsepticProcessing**

Aseptic Processing, Visual Inspection & Packaging *Summit*

Sterile drug product manufacturing is being reshaped by EU GMP Annex 1, advanced barrier technologies, automated visual inspection and increasingly complex primary packaging. Robust contamination control strategies are now expected from filling line to final container.

Over two days in Amsterdam, industry leaders share practical case studies on aseptic processing, visual inspection and packaging integrity, with open Q&A, panel discussions and dedicated networking.

Key Practical Learning Points:

- Applying EU GMP Annex 1 (2022) expectations in daily aseptic operations
- Building and maintaining a practical Contamination Control Strategy (CCS)
- Isolator and RABS operation, decontamination and interventions
- Aseptic Process Simulation (media fill) design and lessons learned
- Manual, semi-automated and automated visual inspection – qualification and defect management
- Container closure integrity and primary/secondary packaging challenges
- Inspection readiness and regulatory expectations for sterile products



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