

LabAnalysis Group is hosting a webinar series to share their knowledge on challenges faced by companies of Pharmaceutical Market on a global scale. Producers stand to gain valuable insights from our webinars for different reasons:

Understanding Regulatory Compliance: we often cover regulatory requirements and guidelines set forth by authorities such as the FDA (Food and Drug Administration), EMA (European Medicines Agency), or other relevant regulatory bodies. Hereby to comply standards and best practices to ensure products meet regulatory requirements.

Ensuring Product Safety and Quality: attending a webinar, producers can learn about the sources, identification, best practices and mitigation strategies for contaminants, thereby safeguarding product safety and consumer health.

Optimizing Product Development: our focus may be on compatibility testing and risk assessment methodologies, empowering producers to make informed decisions and optimize product quality and performance.

Risk Mitigation and Contamination Control: filtration system failures or inadequacies can lead to product contamination, batch rejections, and costly production delays. Our experts provide information on risk mitigation strategies, preventive maintenance practices, and quality assurance measures to minimize the risk of contamination and ensure product sterility throughout the manufacturing process.

Improving Quality Control Practices: we can provide valuable insights into analytical techniques and testing methodologies for identifying and quantifying contaminants. By incorporating these techniques into their quality control processes, producers can enhance product quality, consistency, and compliance with regulatory requirements.

Staying Updated on Industry Trends: our experts will share the latest research findings, regulatory updates, and industry trends, enabling producers to stay ahead of the curve and adapt their practices accordingly.

Join our live webinars for **free**. See details below.

Nitrosamines



09 May • 11:00 - 12:30 (CET)
Language: English

Moderator

Lorenzo Maggi

CEO - LabAnalysis Life Science

Speakers

Claudia Percivalle

Regulatory Affairs Director - AstraZeneca

Alessandro Andolfi

Vice Quality Control - LabAnalysis Life Science

Marco Rossi

Unit HL_IMP Responsible - LabAnalysis Life Science

Erica Tediosi

Study Director GLP Ecotoxicology - LabAnalysis Life Science

Extractables & Leachables



07 November • 11:00 - 12:30 (CET)
Language: English

Moderator

Antonio Legnani

CEO - LabAnalysis Process Pharma

Speakers

Marco Rodda

Scientific Director - Chemsafe Srl

Davide Dova

Unit E&L Responsible - LabAnalysis Life Science

Alessandro Granata

Vice Quality Control - LabAnalysis Life Science

Antonio Legnani

CEO - LabAnalysis Process Pharma

Impact of Annex I on the in-process management of filtration systems in aseptic production and terminal sterilization



13 June • 11:00 - 12:30 (CET)
Language: English

Antonio Legnani

CEO - LabAnalysis Process Pharma

Federica Annovazzi

Technical Manager - LabAnalysis Process Pharma

Case history - Groupe Pemflow

Upon request we can provide a **Certificate of participation** for attending

SUBSCRIBE FOR FREE 

