nuvalore

YOUR PARTNER FOR INNOVATIVE & RELIABLE GMP ANALYSIS

Accelerate your pharma pipeline with confidence. Nuvalore specializes in GMP-compliant analytical testing services for pharmaceuticals. With deep expertise in Analytical Quality by Design (AQbD) and alignment with the latest ICH Q14 guidelines, we deliver data-driven, regulatory-ready results that enable pharmaceutical companies to accelerate every stage of drug development, ensure compliance, mitigate risks, and protect product quality with confidence.

ANALYTICAL SERVICES THAT DELIVER

End-to-end analytical services for biologics, peptides, small molecules, and more.

Analytical Method Development & Optimization

- QbD-driven method development
- DoE-based parameter optimization

Method Development, Validation & Transfer

- Develop robust methods using DoE and conventional tools
- Align with ICH Q14 guidelines
- GMP-compliant reporting

Drug Stability Testing Services

Consulting Services

- ICH-compliant strategies
- GMP application and compliance
- Technology transfer support
- Lifecycle management planning

Drug Characterization & Routine Analytics

- Method Development & Optimization
- Pharmacopoeia Methods
- Peptid-Mapping (PepMap)
- Cation Exchange HPLC (CEX)
- Hydrophobic Interaction
- Chromatography (HIC)
- Hydrophilic Interaction Liquid Chromatography (HILIC)
- Capillary Isoelectric Focusing (cIEF)
- Capillary electrophoresis (CE)
- Characterization of Surfactants (PS80, Pxm188)
- Force Measurement of Prefilled Syringes and Autoinjectors
- Dissolution-Tests
- Measurements of nanoparticles

WHY LEADING PHARMA CHOOSE NUVALORE

Innovation-Driven – Always applying the latest science & technology
Tailored Support – Every project is customized to your exact needs
Trusted Expertise – Decades of experience in pharmaceutical analytics

GET IN TOUCH

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