



Pharma & Nutra services

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Our laboratories, equipped with the most modern instrumentation including UPLC, HPLC, LCMS, LC-MS/MS, LC-TOF-HRMS, GC, GC- and dissolution systems is operated by a well trained and highly motivated staff with an average of over 10 years of practical experience. The Pharmaceutical Analytical Laboratory is GMP inspected and under control of the respective authorities.

We deliver analytical services on:

- Active Pharmaceutical Ingredients (API)
- Raw materials
- Excipients
- Solid formulations
 - Tablets (coated and uncoated)
 - Capsules (hard and soft gelatine capsules)
- Liquid formulations (solutions, suspensions, injectables)
- Semi-solid formulations (gels, creams, ointments, suppositories)

Our services include but are not limited to:

- Analysis according to official monographs (PhEur, USP, AOCs)
- Method development/implementation/optimization & validation
- Stability studies
- Cleaning method development and validation
- Forced degradation studies
- Impurity profiles
- Structural Elucidation of Unknowns
- Reference Standard Characterization
- Solubility studies
- Respons factor studies
- Extractable/Leachables
- Residual solvent analysis
- Batch control
- Release analysis
- Process support
- Container closure integrity testing
- Dissolution and disintegration
- Technology Transfer to QA at Contract Manufacturing partner
- QP- Qualified person service
- Patent Infringement - Analytical support
- FDA Dietary Supplements Consulting



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