INTERTEK PHARMACEUTICAL SERVICES LABORATORY

Whitehouse, New Jersey, USA

intertek
Total Quality. Assured.

Intertek's Pharmaceutical Services laboratory in Whitehouse, NJ is an FDA and DEA registered cGMP compliant facility, specializing in complex and routine testing services to support the development of drug products, formulations, medical devices and drug delivery systems. We have pharmaceutical expertise for various GMP and CMC testing, including method development and validation, stability, extractables and leachables, and elemental impurities.

ELEMENTAL IMPURITIES
United States Pharmacopeia (USP) <232> and <233> elemental impurities testing in drug products including screening and quantification of potentially toxic metal impurities

NITROSAMINE IMPURITY ANALYSIS
Detection and quantification of nitrosamine impurities

ELEMENTAL ANALYSIS & TRACE METALS
Supporting pharmaceutical development and GMP production through elemental composition and impurity quantification testing services

MEDICAL DEVICE TESTING
Analysis to ensure materials comply with strict industry specifications to meet application performance requirements for intended use in industry

STABILITY STUDIES
Supporting drug product development, commercial stability studies, batch release and quality control testing

QUALITY CONTROL
Release testing of raw materials, excipients, APIs and intermediates to pharmacopoeia specifications supported by troubleshooting and specialist QC methods

EXTRACTABLES AND LEACHABLES
Quantification and assessment of risks associated with potential leachable impurities that originate from pharmaceutical container closures, process equipment and medical device packaging

ANALYTICAL METHOD DEVELOPMENT AND VALIDATION
Supporting drug development delivering regulatory-driven and phase-appropriate methodology across a range of analytical technologies

FOR MORE INFORMATION

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