

Contract GMP analytical laboratory services

Pharmaceutical Testing Services Australia

TGA-accredited & US FDA-inspected GMP facility

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Pharmaceutical Testing Services Australia

Trusted GMP Analytical Testing for Pharmaceutical Development, Manufacturing and Quality Control

Intertek's pharmaceutical testing laboratory in Melbourne, Australia, provides GMP analytical testing and scientific support for pharmaceutical, biotechnology and healthcare companies across Australia, New Zealand and global markets.

With more than 35 years of pharmaceutical analytical expertise, our scientists deliver high-quality, regulatory-compliant testing services that support product development, manufacturing, stability programmes, batch release and regulatory submissions. Operating as a TGA and APVMA-accredited and US FDA-inspected facility, we generate reliable data and drive insight of raw materials, actives and finished products, helping clients maintain confidence in product quality.

Why choose Intertek Australia?

- 35+ years of pharmaceutical analytical expertise
- TGA and APVMA-accredited GMP laboratory, with additional APVMA accreditation
- US FDA-inspected facility
- ISO 9001 quality systems
- Advanced analytical instrumentation
- Fast turnaround and responsive technical support
- Part of Intertek's global pharmaceutical laboratory network



Comprehensive GMP testing services

As one of the most capable pharmaceutical chemistry laboratories in Australia, Intertek provides a complete range of TGA compliant pharmaceutical testing services across Australia, New Zealand and Worldwide and is part of the global Intertek Pharmaceutical Services lab network.



Raw Materials Testing

Verification of APIs, excipients and incoming materials to support quality and regulatory compliance.



Method Development & Validation

Development, transfer, verification and validation of robust analytical methods tailored to your products and processes.



Finished Product Testing

Assay, impurities, dissolution, disintegration, residual solvents, elemental impurities and batch release testing.



Manufacturing Investigation Support

Impurity identification, contaminant investigations and troubleshooting to support manufacturing quality and risk mitigation



Stability Studies

Accelerated, long-term, in-use and commercial stability programmes supporting ICH and FDA requirements.



Advanced Analytical Characterisation

Scientific expertise supporting complex analytical challenges and regulatory expectations.

Advanced Analytical Expertise for Pharmaceutical Development

Pharmaceutical products are becoming increasingly complex, creating greater demands for analytical precision, regulatory compliance, and product quality assurance. Intertek Australia combines advanced instrumentation, scientific expertise, and GMP-compliant quality systems to help pharmaceutical and biotechnology companies address analytical challenges throughout the product lifecycle.

Supporting Every Stage of Development

From early-stage development through commercial manufacture, our scientists provide analytical solutions tailored to the evolving needs of pharmaceutical products and processes.



Comprehensive Analytical Testing Capabilities

Our laboratory supports a broad range of pharmaceutical testing requirements, including:

- Assay and related substances
- Residual solvents
- Elemental impurities (ICP-MS)
- Dissolution and disintegration
- ICH Stability studies
- Method development and validation
- Method transfer and verification
- Raw material testing
- Finished product testing
- Impurity profiling and identification
- Contamination investigations
- Physical and physicochemical characterisation
- Classical wet chemistry



Advanced Instrumentation and Scientific Expertise

Our scientists utilise a comprehensive suite of analytical technologies to characterise pharmaceutical materials and products, including HPLC, UPLC, GC, GC-MS, ICP-OES and ICP-MS, FTIR, SEM, NMR, TGA and DSC, dissolution systems, and stability chambers.

By combining advanced instrumentation with practical scientific expertise, we help customers address complex analytical challenges, identify root causes, support regulatory submissions, and maintain confidence in product quality throughout the product lifecycle.

Part of a Global Pharmaceutical Services Network

As part of Intertek Pharmaceutical Services, our Australian laboratory provides local expertise backed by a global network of GMP laboratories, scientists, auditors, and regulatory specialists. This integrated approach gives customers access to specialist capabilities across analytical development, stability studies, extractables and leachables, packaging evaluation, regulatory support, and quality assurance services.

Whether supporting a single analytical challenge or a long-term outsourcing partnership, Intertek helps pharmaceutical and biotechnology companies reduce risk, accelerate development, and bring high-quality products to market with confidence.





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