



International Laboratory Services

Pharmaceutical Division

Pharmaceutical Microbiology Testing

Our purpose built pharmaceutical microbiology laboratories are fully equipped to analyse raw materials, finished products and medical devices in accordance with Pharmacopoeial methodology.

We are fully versed in EP and USP methodologies and we routinely undertake work for companies around the world.

In addition, our routine testing is performed under the scope of our UKAS accreditation, although not exclusively, we are flexible and can work to your preferred methodology.

ILS Pharmaceutical Division is a GMP compliant, MHRA and FDA Inspected Laboratory.

Our testing capabilities include:

- Sterility Testing
- Microbial Limit Testing
- Bacterial Endotoxins (LAL and Kinetic Turbidimetric Assays)
- Antibiotic Assay
- Bioburden Testing
- Preservative Efficacy Testing (PET)
- Water Testing
- Irradiation Dose Setting determination studies to ISO 11137
- Method Development and Validation

Our sterility testing suite conforms to ISO Class 5 Specification.

We have three walk-in incubators and a clean room specifically for endotoxin work.

ILS has a vastly experienced team and can advise on regulatory issues, studies, scientific evaluation of safety and efficacy data and advise on applications for marketing authorisation.

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