

One Solution, One Laboratory

FDAS has an experienced analytical team that come from pharmaceutical backgrounds. This helps us to understand your needs as a manufacturer, the unique pressures you face and your requirements of an analytical laboratory:

- MHRA - audited and approved
- GMP - fully compliant; all samples go through our single QMS
- Expertise - comprehension of the analysis performed and what the results mean to you
- Communication - notification of sample receipt, analysis date, OOS investigations etc
- Single site - chemistry, microbiology, sterility and stability facilities at our Nottingham site
- Absolute confidentiality

Our processes are designed to help you achieve maximum efficiencies by being as flexible as possible. Only in this way can we become an effective and integral part of your QMS.

For example you can:

- Pre-book test analysis slots
- Receive specifically formatted documentation
- Utilise flexible payment structures
- Have a rapid response to quotations

Quality Assured, Service Guaranteed



Sample Analysis

Analytical Development and Validation

Speciality Chemicals

QC Analytical Services

Stability Studies



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