Validating Your Cleanroom Facility

ENVAIR TECHNOLOGY

Clean Air & Containment

Envair Technology provide comprehensive commissioning and validation services for cleanroom facilities, verifying compliance with ISO 14644 and Good Manufacturing Practices (GMP) standards.

From large-scale pharmaceutical production facilities to smaller pharmacies, our team of highly trained engineers will ensure your equipment maintains its integrity, efficiency and compliance.

With over 50 years of excellence and innovation, we have built a trusted reputation for servicing all brands of clean air and containment equipment. Contact us today for a quick, efficient solution that minimises downtime and maximises the lifespan of your equipment.

COMPREHENSIVE CLEANROOM SUPPORT

At Envair Technology, we don't just design, manufacture and install cleanroom products – we also keep your facility running at peak performance too. We offer ongoing support for any brand of containment equipment, including:

- Servicing and maintenance
- Validation and testing
- Genuine spare parts and consumables
- Routine and emergency VHP fumigation

To schedule your validation, request urgent assistance or find out more, please contact us:

01706 228 416
service@envairtechnology.com

CLEANROOM VALIDATION SERVICES

Our cleanroom validation service includes a wide range of tests to ensure compliance with all the latest ISO standards and industry regulations.

Routine Tests:

- Airflow measurements
- Pressure regimes
- HEPA filter integrity testing
- Particle counting

Additional Tests:

- Airflow visualisation studies
- Recovery rate analysis
- Microbiological sampling

PROTOCOL WRITING AND COMPLIANCE DOCUMENTATION

Alongside our practical validation services, we offer protocol writing tailored to your specific needs, ensuring full compliance with the regulations of your industry.

Our documentation services include:

- User Requirement Specification (URS)
- Installation Qualification (IQ), Operational Qualification (OQ) and Performance Qualification (PQ) documentation
- Onsite testing and compliance checks
- Certified results of all compliance verification tests: delivered electronically within agreed timescales

We are BS EN ISO 9001 certified and always work to the highest quality standards. With decades of experience in cleanroom validation and containment solutions, our UK wide team of expert engineers are ready to support your facility.

bsi ISO 9001 Quality Management Systems CERTIFIED

Whatever you need to maintain the validation of your cleanroom, Envair Technology have you covered with our comprehensive range of support services.

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QUALITY STATEMENT

Envair Technology provide world class design and manufacturing expertise for a full portfolio of cleanroom and containment solutions. We work with healthcare, pharmaceutical and laboratory facilities around the world, to protect the safety of teams handling hazardous compounds and maintain product integrity. Our engineers and production team are committed to the highest standards of excellence, meeting the requirements of ISO 9001:2015. We provide a comprehensive maintenance, testing and parts service for all our products, to ensure a lifetime of trouble-free use.

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