*symbiosis
Pharmaceutical Services



Your Aseptic Injectable CDMO

Symbiosis manufacturing facilities are FDA-inspected and MHRA-licensed and were purpose-built to support the GMP fill-finish manufacture of injectable drug products for clinical trials and commercial sale globally.

Offering Annex 1 compliant cleanrooms (including PUPSIT), alongside BSL-2 and cytotoxic capabilities, automated filling and visual inspection, Symbiosis manufactures to the most stringent current industry standards.

Complemented by our in-house QC Laboratories, we can support the manufacture of your drug product from development to QP release.

Our Aseptic Manufacturing Capabilities

Symbiosis Services Include:

- Bulk formulation / Sterile Filtration
- Aseptic GMP Fill-finish manufacture
- Analytical and Microbial Testing
- Labelling and Packaging
- QP Release
- GMP Storage / Distribution



Our State of the Art Facilities offer:

- Up to 15,000 vials per batch
- 200 L bulk volume
- Automated visual inspection
- Liquid or lyophilised formulation

Ready to explore our facilities?

Discover how we can accelerate your drug manufacturing timelines

THE KING'S AWARDS FOR ENTERPRISE 2025 INTERNATIONAL TRADE

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