



GSK Contract Manufacturing Biopharmaceuticals



GlaxoSmithKline Biopharmaceuticals is a full-service contract manufacturing partner in the development and manufacture of cGMP biopharmaceutical proteins for clinical trials and commercial use. We offer FDA and European licensed multi-product facilities. Our featured services include the low-cost, rapid scale-up and manufacture of mammalian cell-derived proteins at up to 5,000L scale and the preparation of formulated bulk products.

We also offer fill/finish for sterile liquids and freeze-dried drug products. Our offerings include complete analytical and regulatory services to speed technical transfer and regulatory approval. GSK is at the forefront of process optimization with the application of methodologies that include Process Analytical Technology (PAT) and Quality by Design (QbD). Our manufacturing has support from comprehensive corporate resources, including Chemistry and Pharmaceutical Development.

Our independent product purification suites provide complete segregation of a client's product. We offer state-of-the-art testing laboratories with a full array of testing capabilities, advanced computer tracking and control, and cGMP compliant warehousing for raw materials and formulated bulk product.

GlaxoSmithKline Biopharmaceuticals is expanding the proud heritage of discovery and development established by our parent company, GlaxoSmithKline. With bulk manufacturing sites (US), secondary filling (UK and Italy), and a central testing lab (UK), we have the experience, knowledge, resources, and corporate backing to satisfy all requirements for reliable, cost-effective, and high-quality biopharmaceutical contract manufacturing services

Molecule to Market — Extending Your Capabilities

Manufacturing Capabilities

Large Molecule / Biopharm

- Large company strength and small company flexibility
- Cost-effective operational scale
 - 75L to 5000L bioreactors in three manufacturing suites
 - 6 production trains with three purification suites
- Over 10 years with a successful track record in transferring and manufacturing multiple products
- Ability to work with a broad range of clients including major pharmaceutical companies
- Highly developed and proven methods for technology transfer
- Fully trained staff experienced in cGMP manufacturing, testing, and release
- Complete supply solution (bulk drug substance manufacture, fill/finish, European release)

- Proven executional excellence: on-time, on-spec performance
- Protection of proprietary technology
- Exemplary and sustained safety record

Processes and Equipment

- Independent Cell Culture Suites — producing product up to 5,000L scale, with multiple 5,000L scale bioreactors
- Independent Product Purification Suites — providing complete segregation of a client's product
- State-of-the-Art Testing Laboratories — with a full array of testing capabilities
- Advanced computer tracking and control
- cGMP Compliant Warehousing — for raw materials and formulated bulk product

