



AmbioPharm Inc

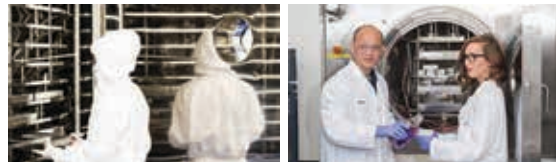
ACCELERATING YOUR PEPTIDE TO MARKET

Peptide Contract Manufacturing & Development Services

Company Profile

ACCELERATING YOUR PEPTIDE TO MARKET

AmbioPharm (APi) is a global, full-service, contract development & manufacturing organization (CDMO), with a focus on the production of peptides. Our facilities include our headquarters and USA campus located in North Augusta, South Carolina, as well as our China campus located in Shanghai. Our services include process development and optimization, manufacturing of building blocks and key raw materials, and custom peptide synthesis. We are capable of manufacturing from mg to multi-kg scale to produce bulk peptides to custom specifications, using a wide range of skill-sets in both solid-phase, solution-phase, or hybrid chemistry approaches. We also perform organic conjugations to small molecules, proteins, toxoids, antifungals, KLH, and PEG. With a leadership team of peptide experts consisting of over 150 years of cumulative experience running pilot to commercial scale production, AmbioPharm is ready to support all of your peptide needs.





South Carolina, US Campus (Headquarters)

More than 250 employees staff the USA campus where a huge expansion project was completed in 2020, adding 55,000 square feet of space to the existing campus, totaling 82,000 square feet. The new buildings include suites to increase our capacity for purification, lyophilization, analytical QC testing, and stability and warehouse storage.

Shanghai, China Campus

In Shanghai, China, more than 375 employees on this campus are supporting our peptide manufacturing and testing operations. An impressive 300,000 square foot state-of-the-art manufacturing building in Shanghai has been constructed, growing our total campus footprint to 380,550 square feet. This expansion allows us to more than double our synthetic chemistry capacity, and support the growth from current and new customers in both small and large scale peptide production.

Non-GMP & Development Services

SUPPORTING YOUR R&D NEEDS

Key R&D Equipment & Capacity

CEM Liberty Blue HT-12 Automated Microwave Peptide Synthesizers

(up to 12 sequential peptides; rapid HTP work)

Manual Glass Reactors (100, 150, 250, 500mL, & 1L)

Multiple Large Scale 2, 5, 10, & 20L SPPS Reactors

Key R&D Analytical & Purification Equipment

Shimadzu and Agilent Analytical HPLC Systems

Small & Large Scale Purification Systems

(multiple HPLC self-pack axial pressure columns: 5, 8, 15, & 20cm)

SG100 Protein Purification Systems

HPLC automated Semi-Prep HPLC Systems

Production & Analytical MS & HPLC Systems



Process Development Programs

- Initial synthesis usually in Fmoc-tBu format for SPPS
- Hybrid methods if scale or length warrant this approach
- Solution phase approach for large volumes projects
- Scouting synthesis run in a fully automated approach using conventional and microwave techniques
 - LC-MS analysis of crude product to determine efficiency of synthesis and where any failures appear
 - Resynthesis with special resins (e.g. TentaGel, Chem-matrix, CTC, etc.) with substitutional variation
 - Resynthesis also with special building blocks such as Psuedopro to minimize resin aggregation and eliminate tough deletion peptides (Gly, Ser, Pro, Thr)
 - Compile data and determine optimal choice of resin, substitution, building blocks and protecting groups
 - Initiate secondary steps such as cleavage, oxidation/folding, purification optimization
- Perform SPPS with the above selections and analyze crude product (automated)
- Begin scale-up with optimized synthesis protocol (usually manual)
- Finalize process optimization for cleavage & oxidation, cyclization etc.
- Develop preparative HPLC conditions for purification and analysis
- Scale-up HPLC purification
- Establish analytical RP-HPLC method for purity
- Establish salt-exchange and lyophilization conditions
- QC specifications: determine & report
- Technical document preparation



Our Offerings

GMP Peptides

AmbioPharm has multiple manufacturing suites capable of producing peptide APIs (Active Pharmaceutical Ingredients) at varying scales from gram to multi-kilogram (> 100 kg) scale. We use solid phase synthesis, solution phase synthesis, hybrid solid and solution phase synthesis, and native chemical ligation to produce peptide APIs. Through a partnership with CEM Inc., we can also perform GMP production using their patented, large-scale Liberty Pro™ microwave peptide synthesizers in our in North Augusta, SC facility.



Non-GMP Peptides

In addition to producing peptide products under GMP standards and requirements, we also provide process development and analytical services for non-GMP grade peptides. We have added automated peptide synthesizers to help speed our delivery of these non-GMP research peptides. These services are available for a broad range of customers, including universities, research organizations, biotechnology, pharmaceutical, cosmetic, agricultural, veterinary companies, and beyond.

Generic Peptides

AmbioPharm is one of the largest capacity peptide API manufacturing companies in the world. With a USA campus and headquarters located in North Augusta, SC and facilities in Shanghai, China, we manufacture both new chemical entities and generic peptides under GMP for clients worldwide. We also partner with generic drug companies to develop generic peptide drugs (both API and Drug Product) for established commercial markets worldwide.



Analytical Development & Validation

As your clinical or commercial programs require, AmbioPharm has established analytical teams with specific peptide expertise to support the development and phase appropriate validation of methods. These methods allow us to monitor the process for side reactions, meet required purity standards, and limit any residual impurities in the product. Controlling the final API to ensure the required quality attributes are achieved, while minimizing process and degradation impurities, affords formulators the maximum labeled drug shelf-life under proper storage conditions.

Vaccine Services

AmbioPharm has years of experience manufacturing Active Pharmaceutical Ingredients (APIs) for peptide vaccines in varying configurations. Our experience in high-quality manufacturing allows us to make the peptide construct within GMP standards and provide both quality and value. We also provide guidance on expertise in three key areas that are critical to successful implementation: analytical, formulation, and regulatory.

Our Capacity

Synthesis Capacity

Solid Phase Reactors:

(standard or microwave*)

30mL*, 1, 3, 5, 8*, 10, 15*, 30, 50, 80,
200, 500, 1000, & 3000L
(up to 100kg crude/batch)

Solution Phase Reactors:

10, 20, 30, 50, 80, 100, 200, 300, 500,
1000, 1500, 2000, 3000, & 5000L
(up to 500kg crude/batch)

Hybrid Synthesis

(up to 500kg crude/batch)



Lyophilization Scale

Multiple Manifold Lyophilizers
(up to 1kg/batch and 50kg/year)

100, 200L Tray Lyophilizers
(up to 6kg/batch and 250kg/year)

400, 500L Tray Lyophilizers
(up to 8kg/batch and 400kg/year)

800, 1000L Tray Lyophilizers
(up to 15kg/batch and 800kg/year)

Purification Scale

Automated State-of-the-Art HPLC Large Scale Purification

Preparative HPLC columns:

ID: 5, 8, 15, 20 & 30cm HPLC Columns
(up to 7kg/batch and 100kg/year)

ID: 45cm HPLC Columns
(up to 15kg/batch and 120kg/year)

ID: 60cm HPLC Columns
(up to 28kg/batch and 200kg/year)

ID: 100cm HPLC Column
(up to 40kg/batch)



Quality & Compliance

AmbioPharm maintains the highest level of quality and compliance in its manufacturing plants in North Augusta, SC, and Shanghai, China. As one of the leading suppliers of peptide APIs worldwide, we provide rigid quality control, SOPs, training, and rigorous testing.

Key analytical testing which we offer includes: HPLC, UPLC, GC, LC-MS, IC, SEC, KF, Nitrogen Content, AAA, N-terminus Sequencer, Endotoxin, Bioburden, and more.

Why Us

Capability



Wide range of technologies, equipment, & scale.

Experience



Extensive peptide & GMP management expertise.

Capacity



Industry leading manufacturing capacity & expansion plans.

Globalization



Multiple manufacturing sites & global customer support.

Quality



Systems & infrastructure to support commercialization of peptides.

Speed




Resources & infrastructure to ensure shortened delivery times.



AmbioPharm, Inc.

Accelerating Your Peptide to Market

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