



BRINGING
YOUR
SCIENCE
TO LIFE

Bioconjugation and
Antibody Drug Conjugates



Your Partner for Conjugation Projects



- **Pioneers in the highly potent landscape for over 14 years, we successfully managed numerous drug-linker projects. Since our first ADC project in 2005, many customers, ranging from small biotech to large pharmaceutical companies, expressed a growing interest in our ADC and bioconjugation abilities. We can manage projects from drug-linker to final drug product.**

CARBOGEN AMCIS offers tailored programs based on your needs and budget. You will benefit from our dedicated team of experienced chemists, biochemists and biologists, who can advise you on the best linker and conjugation strategy. For all other projects conducted by CARBOGEN AMCIS, safety and quality are central in the ADC area.

“We advise our customers on the best conjugation strategy to successfully manage projects from drug-linker synthesis to final drug product manufacture.”

Luca – Senior Chemist ADC

Why choose us?

- 1 Project management experience, dedicated team, seamless and personalized communication with customers
- 2 cGMP process development and manufacture of warheads and linkers (OEL < 0.01 ug/m³)
- 3 cGMP conjugation of mAb to small molecules and polymers
- 4 Clean room suites fully qualified for cGMP manufacturing dedicated to bio-conjugation
- 5 State of art purification technologies
- 6 Outstanding analytical capabilities
- 7 Fill-and-finish capabilities
- 8 Regulatory and CMC support

Conjugation & Purification Capabilities

- Manipulation of biological molecules
- Chemical modification with linkers and activators
- Conjugation of monoclonal antibodies and biologics to small molecule compounds up to 20 L volume or ca.300g ADC per batch
- ÄKTA-Pure 25 chromatography system
- ÄKTA Ready Skid for large scale aseptic
- PALL SU-TFF up to 2.5m² automated system for aseptic diafiltration
- OEL 10 ng/m³ 8h-TWA
- Aseptic environment with Grade D and C qualified areas
- cGMP compliant
- Single use liquid handling systems



Grade D Clean Room, Bubendorf, Switzerland

Analytical & Bioanalytical Capabilities

- In-process monitoring by various means such as: UV/Vis, HPLC, Endo-Safe, pH and conductivity
- ICH stability studies
- Full on-site analytical support for the release testing of the product
- Enzyme Linked Immunosorbent Assay (ELISA)
- SDS-PAGE (Gel Electrophoresis)
- iCE3 Imaged Capillary Isoelectric Focusing
- Bioanalyzer Agilent 2100
- Endotoxin analysis
- HPLC analysis (RP, SEC, HIC, PLRP)

Fill & Finish Capabilities

In addition to our process research and manufacturing services for the fast supply of highly-potent APIs, CARBOGEN AMCIS offers formulation services for highly-potent drug products and cGMP aseptic production of parenteral drugs, including cytostatics and cytotoxics..

- Pre-clinical batches (for technical, stability or toxicology studies)
- Clinical batches (for phases I, II and III)
- Injectables: liquid and freeze-dried forms



Sterile Fill & Finish Room, France



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CARBOGEN AMCIS AG is a leading service provider, offering a portfolio of drug-development and commercialization services to the pharmaceutical and biopharmaceutical industry at all stages of drug development. The integrated services provide innovative chemistry solutions to support timely and safe drug development allowing customers to make the best use of available resources.

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