CARBOGEN AMCIS

A Dishman Group Company

BRINGING YOUR SCIENCE TO LIFE

Bioconjugation and Antibody Drug Conjugates

Your Partner for Conjugation Projects

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Pioneers in the highly potent landscape for decades, 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.

CARBOGEN AMCIS specializes in ADC strategies, providing expertise in every aspect.
Watch our video.

Why choose us?

Drug-linker
cGMP process development and manufacture of Warheads & Linkers (OEL < 0.01 µg/m³)

Purification
State of art purification technologies

Conjugation
cGMP Conjugation of mAb to Small

Dedicated Team
A multi-disciplinary and extensively experienced team

Analytical
Outstanding analytical capabilities

Fill & Finish
Fill-&-Finish capabilities

Clean Room
Clean Room fully qualified for cGMP manufacturing dedicated to bio-conjugation

Support
Regulatory and CMC support

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

Luca – Manager ADC

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 2.5m^2 automated system for aseptic diafiltration
- OEL 10 ng/m^3 8h-TWA
- Aseptic environment with Grade D and C qualified areas
- cGMP compliant
- Single use liquid handling systems

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)

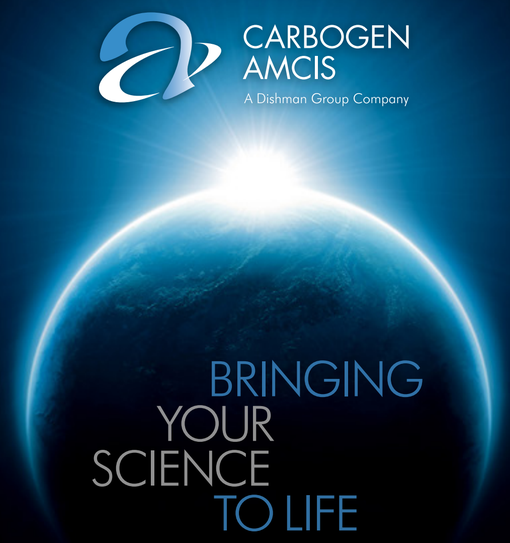
Grade D Clean Room, Bubendorf, Switzerland

Fill & Finish Capabilities

In addition to our process research and manufacturing services for the fast supply of highly potent APIs, we offer formulation services for highly potent drug products and cGMP sterile 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)
- Commercial supply
- Injectables: liquid and freeze-dried forms

Grade C Conjugation Suite, Bubendorf, Switzerland



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