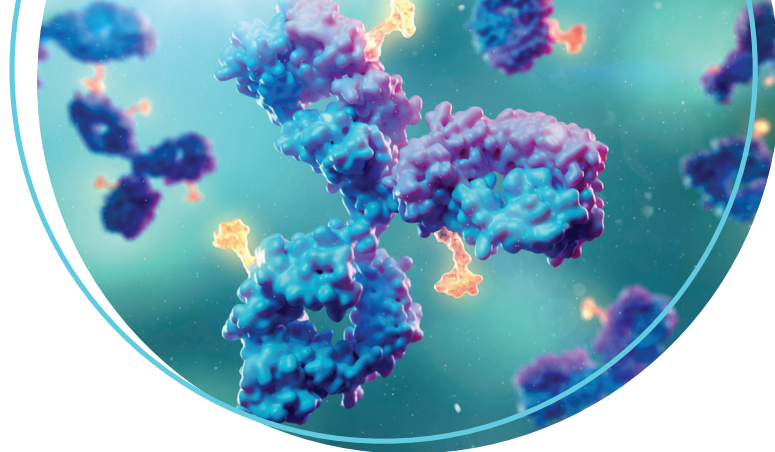




LABCORP BIOPHARMA LABORATORY SERVICES

Antibody-drug conjugate (ADC) capabilities





Delivering global ADC development solutions

Antibody-drug conjugates (ADCs) continue to be a widely used and key modality of interest for the pharmaceutical industry due to their ability to deliver targeted cytotoxic payloads to solid tumors and blood-based malignancies that induce cell apoptosis for oncology patients.

While payloads and antibody types continue to evolve, ADCs remain challenging given their highly complex and nuanced chemistry. For an ADC to be successful, the right combination of target-antigen, active linker payload, appropriate drug-to-antibody ratio (DAR), proper tumor indication, and payload conjugation site need to be considered.

Given this complexity, ADC drug developers face specific challenges in bringing their treatment candidates through the drug development continuum, including:



Toxicity

Elevated risk with high DARs and differing payload conjugation configurations



Immunogenicity

ADC antibody, linker, and payloads can have different impacts on the immune system response



PK/PD

More complex and frequent assessments given antibody, linker, and payload MOAs



Payload potency and efficacy

Tumor cell apoptosis and continued efficacy over time



Off-target effects

Unintended negative impacts beyond the tumor



ADC characterization and stability

Demonstrating DAR assessments and manufacturing process stability



Biomarker and companion diagnostic (CDx) development

Patient stratification and treatment for clinical trials as well as Day-1 post-approval readiness

How Labcorp Biopharma Laboratory Services can help

Labcorp Biopharma Laboratory Services is a highly experienced, comprehensive, and global ADC development partner with laboratory capabilities spanning early development through commercialization as well as expertise in oncology biomarkers and CDx co-development capabilities.



Supported 15 of the 18 **(83%) approved ADCs** currently on the global market



Is supporting 13 of the 17 **(77%) mid to late clinical trial ADCs** currently in global development



Has experience **with multiple generations of payloads**, including DNA-damaging agents, tubulin binders, and topoisomerase 1 inhibitors, as well as newer payload MOAs

Our specific ADC drug development and biomarker/CDx capabilities



Discovery

- Exploratory/nonregulated ADC DMPK, safety, *in vitro* assessments
- *In vivo* pharmacology, imaging, and focal radiation capabilities
- Human and syngeneic tumor models available



Toxicology/Immunotoxicology/Immunogenicity

- Nonclinical biodistribution and toxicokinetic ADC impact on relevant animal models (primate homologue, rodents for novel linkers and toxins) and human tissues
- Tumor samples to assess drug candidate targeting impact effectiveness
- ADC assessment of antibody, linker, and payloads



Antibody Reagents and Vaccines

- ADA positive control antibodies for ADCs for ADA/PK assays
- Antibody reagents for ADCs



Bioanalytical Services

- GLP/GCP nonclinical and clinical ADC PK/PD via LBA/LC-MS
- Quantity of free toxin, total antibody (+DAR 0), and drug-conjugated antibody (DAR > 0)
- Immunogenicity via ADA/nAb assessments



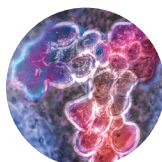
Companion Diagnostics (CDx) Development

- CDx strategy consultation development, including design control, assay development, analytical and clinical validation, and commercialization services*



Central Laboratory Services

- Safety testing
- Tumor assessment (immunohistochemistry)
- ADC receptor occupancy (flow cytometry)
- Novel methods to monitor internalization/payload release
- Biomarker strategies and CDx support within clinical trial environment
- Immunotherapy (IO) and gene expressions/mutations



Chemistry, Manufacturing and Controls

- ADC characterization
- Integrity of antibody and payload conjugation/location
- Bespoke methods for ADCs
- ADC QC release**

*IVD being used for patient management, invasive sampling beyond standard of care, and/or high risk to patients

**QC release services available for U.S. and U.K. locations only; neat toxin reference standard lab safety limitations – requires discussion

Need assistance advancing your ADC drug development program?

Labcorp Biopharma Laboratory Services can help. Please visit our website to submit a contact request or contact your local sales associate for further information.

Visit us at

labcorp.com/modalities/vaccines/antidrug-conjugates



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