GMP Manufacturing Services

Utilizing custom design and development to achieve regulatory approval

ABL has a long and successful history as a CDMO providing contract development and GMP biomanufacturing services to the biotech and biopharmaceutical industry. Our primary focus is viral vector manufacturing and we operate GMP facilities in the U.S. and Europe to meet your global regulatory and redundancy requirements. ABL offers a comprehensive portfolio of effective design, development, production and testing services for a broad range of virus products.

The success of your novel product starts with ABL

ABL’s 30+ years of viral-vector development and GMP production accelerates your novel gene therapies, vaccines, oncolytics and immunotherapies.

30+

YEARS OF EXPERIENCE

200+

CLIENTS

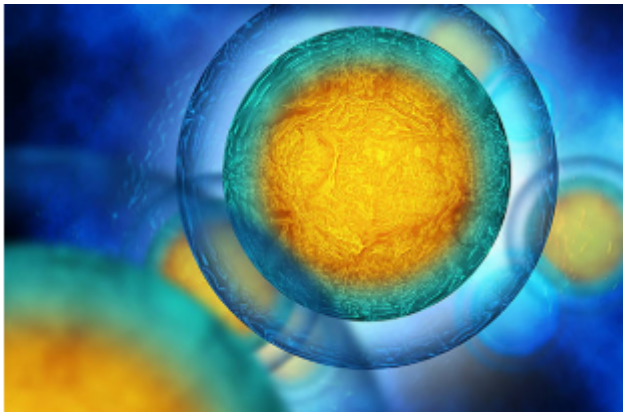
250+

GMP LOTS

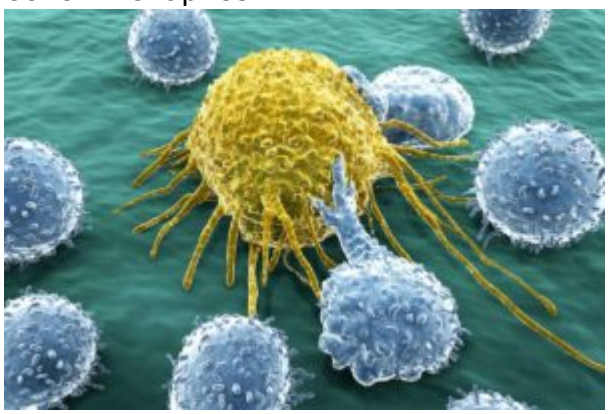
Viral Vector Manufacturing Experts

Our product types

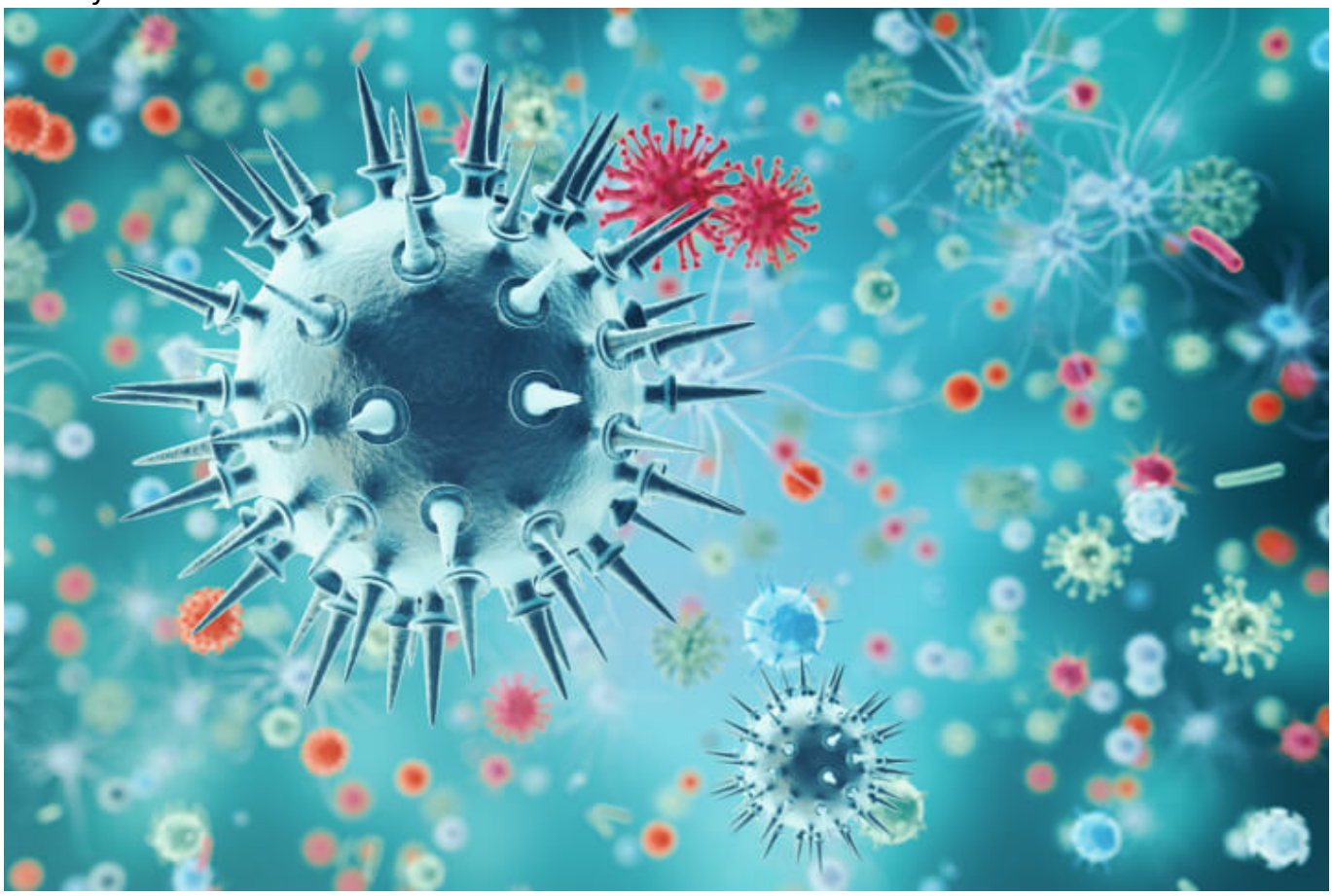
We are a biomanufacturing corporation with global facilities, suites and production platforms focused on accelerating your life changing innovation to market



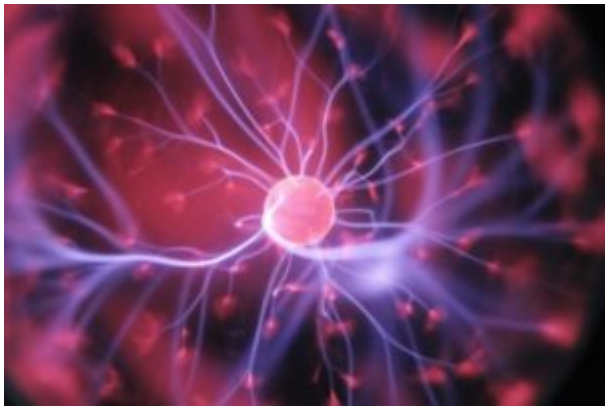
Gene Therapies



Oncolytics



Vaccines



Immunotherapies

Experience producing numerous viral vectors for

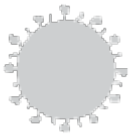
Gene therapies | Vaccines | Oncolytics | Immunotherapies

HSV-1

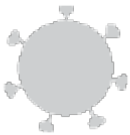
Vaccinia (and MVA)

VLPs

Zika



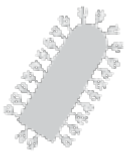
CMV



Lentivirus
(and Retrovirus)

Adenovirus

AAV



VSV

When developing life changing innovations,
the right strategic partner makes the difference

Comprehensive GMP Manufacturing Support

Process design

Process feasibility

Upstream and/or downstream process development

Assay development and qualification

Master and working cell and virus banks

Scale-up and engineering run(s)

GMP bulk drug substance

GMP aseptic fill/finish

Analytical lot release testing

ICH stability testing

Chemistry, manufactoring and controls support

Quality systems

ABL’s U.S. location adheres to Good Manufacturing Practice (GMP) standards and is ISO 9001 accredited. ABL Europe’s facilities are regularly inspected by the French regulatory authorities (ANSM) and is a GMP licensed site for the manufacture of drug substance and aseptically prepared small volume liquid (drug product) viral products in accordance with EMA regulations.

Global GMP Manufacturing Facility

Our U.S. facility is available for GMP production of clinical, pre-clinical trials, and commercial material with iCELLis 500+ platform ready for use. See our New Capabilities.

Rockville, MD, USA

Learn about our biomanufacturing services

- Viral manufacturing
- Process development
- Aseptic fill/finish
- Biologics and QC testing
- Oncolytics

In addition to the services listed above, ask about our protein manufacturing and GMP antibody production services.

Specialist ready to execute. Let’s start the discussion today.

contact us today