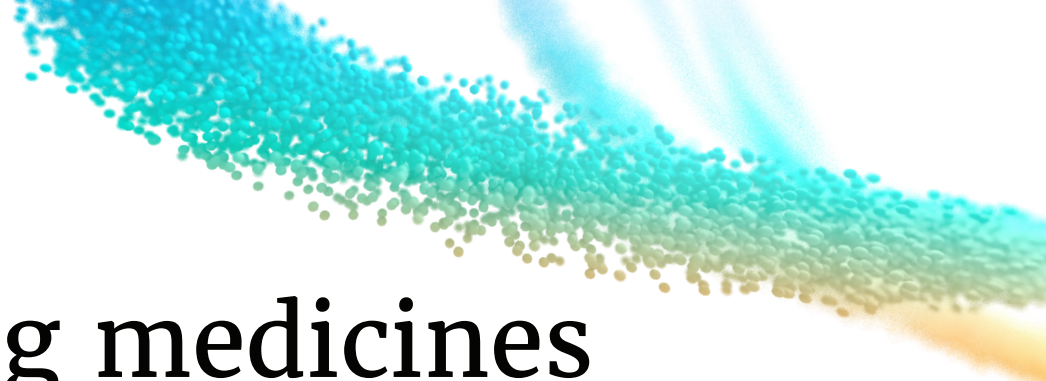




[RESILIENCE

Advancing medicines that *matter*.

Built on Experience. Focused on What's Next.

Resilience is the CDMO partner for biotech and pharma companies at every stage of development and manufacturing.

Our teams bring more than 20 years of expertise across biologics drug substance and sterile drug product, with device assembly, labeling, and packaging capabilities that extend through market-ready finished goods.

Across our North American network, experienced teams, advanced technology, and proven manufacturing operations turn promising science into dependable supply. And from day one, we plan for what comes next, giving every program the best chance to become a medicine that matters.

220M+

Sterile drug
product units
released

600+

Batches
executed
each year

50+

Global
markets
served

50L–2,000L

Single-use bioreactors

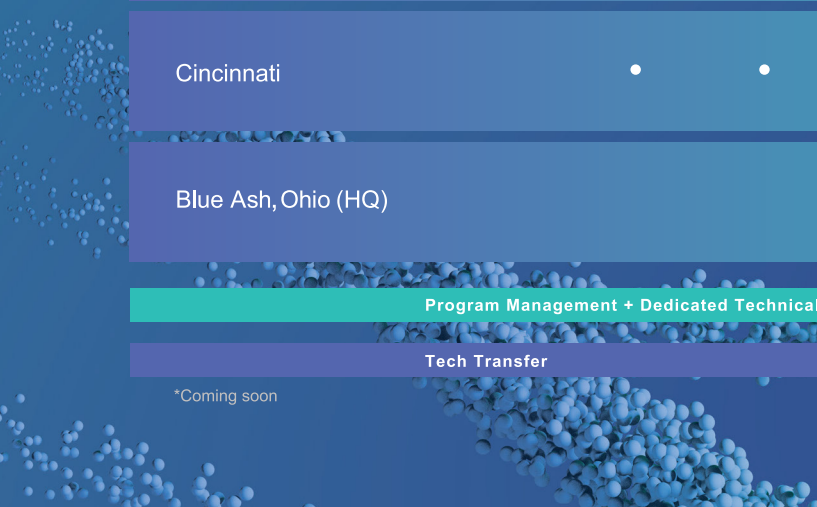
1M+

Square feet of
manufacturing
facilities in North
America

Where – *and how* – we can advance your program.

	Pre-Clinical				Clinical			Commercial	
Sites	Cell Line Development	Process Analytical Development	Scale-up & Optimization	Testing & Stability	Drug Substance	Formulation Development	Drug Product	Device Assembly, Packaging & Logistics	Commercially Licensed
Toronto	•	•	•	•	50L - 2,000L	•	Vial Prefilled Syringe*		YES
Cincinnati			•	•		•	Vial Prefilled Syringe Cartridges	•	YES
Blue Ash, Ohio (HQ)								•	2027*
Program Management + Dedicated Technical Lead									
Tech Transfer									

*Coming soon



Capabilities for Clinical and Commercial Programs

Start at any phase of your program. Choose drug substance, drug product, or both - whether that means beginning with development and tech transfer, scaling up later, or bringing us material you've already made.

Development

Process, analytical, formulation development, and process characterization.

Biologics Drug Substance

Monoclonal antibodies, recombinant proteins, and other complex molecules as well as plasma proteins and transgenics.

Sterile Drug Product

Formulation and aseptic filling of vials, prefilled syringes, and cartridges.

Device Assembly, Packaging & Logistics

Autoinjector and prefilled syringe assembly, multiple vial packaging options, plus labeling, serialization, and cold-chain logistics.

Specialized North American Operations

Blue Ash Greater Cincinnati, Ohio Area

Headquarters

- Automated visual inspection
- Device assembly, labeling, packaging
- Cold storage and logistics
- Driven by advanced equipment and dedicated operational teams

West Chester Greater Cincinnati, Ohio Area

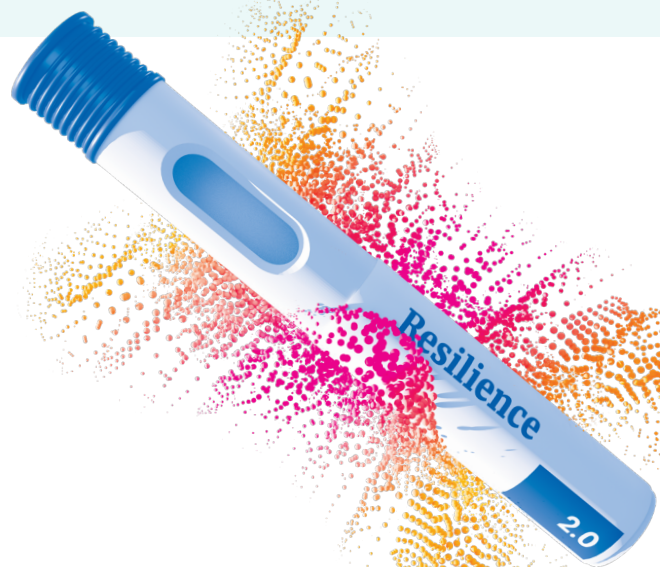
Sterile Drug Product

- Aseptic fill/finish manufacturing
- Automated visual inspection
- Device assembly, labeling, and packaging
- Enabled by digital operations and established sustainability initiatives

Toronto Ontario, Canada

Biologics Hub

- Process and analytical development
- Formulation
- Aseptic fill/finish
- Cold storage
- Comprehensive logistics



Safety First, Quality Always.

- 100% regulatory inspection success rate across our network for more than a decade.
- 100% pass rate for third-party and EU Annex 1 assessments.
- Technical depth and demonstrated expertise across process and analytical development, formulation, process characterization, technology transfer, and validation.

Successful inspections

- USFDA cGMP
- USFDA PAI
- USFDA PFS & Vial Approval
- Health Canada
- EMA Virtual Inspection

More About Resilience

Resilience is a North American contract development and manufacturing organization (CDMO) focused on delivering high-quality, scalable manufacturing solutions for advanced therapies. With all manufacturing in North America, Resilience provides integrated capabilities spanning biologics drug substance, sterile drug product manufacturing, and device assembly and packaging for both small and large molecules. The company partners with leading pharmaceutical and biotech companies to bring complex medicines to market faster and more reliably.



resilience.com
(888) 737-2460