

Summary of capabilities

Lexington, Massachusetts

Facility facts:

Workforce: 320
 Contact Info: 45 Hartwell Ave
 Lexington, Massachusetts 02421
 Tel: 781-266-0200

Unique offering:

63,000-square-foot facility with robust manufacturing capacity in support of high demand for gene therapy products.

Specialized capabilities:

- Process characterization and qualification
- Analytical method validation
- Pivotal and commercial viral vector supply
- State-of-the-art commercial aseptic fill and finish
- Experienced in viral vector manufacturing for cell and gene therapies
- State-of-the-art commercial aseptic fill and finish
- AAV, AdV, RV, HSV, and LV
- Broad array of manufacturing platforms
 - Suspension: Commercial, up to 1,000L scale
 - Adherent: 48HS36 iCELLis® 500

Viral vector services capabilities overview:

Facility offering	Specifications
GMP Manufacturing	<ul style="list-style-type: none"> • Eight drug substance suites and one drug product suite • GMP manufacturing support area • Client reserved capacity • Flexible facility design with all viral suites capable of supporting a range of different technologies including iCELLis® 500, cell factories (flat stock), and up to 1000L stirred-tank bioreactors • Single-pass air HVAC in manufacturing suites
Laboratory Space	<ul style="list-style-type: none"> • QC labs – micro, in-process, release • Manufacturing sciences lab for process establishment
Warehouse Space	<ul style="list-style-type: none"> • Short-term GMP storage space

ADENOVIRAL

Adherent + Suspension
Mammalian cells infection

AAV

Producer cell line + Ad	Adherent + Suspension	Suspension + HSV	Suspension + Baculovirus
Mammalian cells + infection	Mammalian cells transient transfection	Mammalian cells	Insect cells – infection

LENTIVIRAL

Packaging/ Producer cell line	Adherent + Suspension
Mammalian cells	Mammalian cells transient transfection

RETROVIRAL

Packaging/ Producer cell line	Adherent + Suspension
Mammalian cells	Mammalian cells transient transfection

HERPESVIRAL

Adherent + Suspension
Mammalian cells infection

*For detailed capabilities and capacity information please contact your Thermo Fisher Scientific representative.



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From molecule to medicine: An integrated partner for every step in your drug development journey.

Thermo Fisher Scientific provides industry-leading pharma services solutions for drug development, clinical trial logistics, and commercial manufacturing to customers through our Patheon brand. With more than 60 locations around the world, we provide integrated, end-to-end capabilities across all phases of development, including API, biologics, viral vectors, cGMP plasmids, formulation, clinical trial solutions, logistics services, and commercial manufacturing, and packaging. Built on a reputation for scientific and technical excellence, we provide pharma and biotech companies of all sizes instant access to a global network of facilities and experts across the Americas, Europe, Asia, and Australia. We offer integrated drug development and clinical services tailored to fit your drug development journey through our Patheon™ Quick to Care™ program. Our Patheon™ Quick to Clinic™ programs for large and small molecules help you balance speed and risk during early development so you can file your Investigational New Drug Application (IND) quickly and successfully. Digital innovations such as our mysupply Platform and Pharma 4.0 enablement offer real-time data and a streamlined experience. Together with our customers, we're rapidly turning pharmaceutical possibilities into realities.

