

## Summary of capabilities

### Mexico City, Mexico

#### Overview:

Our Mexico facility and operations team offers specialized services in clinical supply chain management and supports global pharmaceutical and biotech companies wishing to conduct clinical research in Mexico and Latin America. Our highly trained project management team has a track record of excellence in the management of clinical supply chain projects. Our multi-lingual team is proficient in helping global pharmaceutical and biotech companies distribute shipments to investigator sites where the majority of shipments require temperature management and temperature monitoring.

Facility facts:		Capabilities:
Opened:	2008	<ul style="list-style-type: none"> <li>• Importation/exportation services including tax and duty payments and customs clearance</li> <li>• GMP storage at ambient, controlled ambient (15°C to 25°C), refrigerated (2°C to 8°C), and frozen (options ranging from -15°C to -80°C) temperatures, as well as ambient and refrigerated controlled drug storage</li> <li>• Secondary packaging in environments including refrigerated (2°C to 8°C) and frozen (-20°C)</li> <li>• In-house labeling, including cold chain and expiry date, labeling/re-labeling</li> <li>• Comparator and clinical ancillaries sourcing and management</li> <li>• Clinical supply optimization services</li> <li>• Pick and pack and distribution services that include temperature management and monitoring for local, regional, and international shipments</li> <li>• Clinical supply returns/storage/destruction service</li> </ul>
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#### Mexico City clinical storage and packaging capabilities:

Storage capacity		Capabilities and services
Ambient (packaging)	1,013 sq. ft. (94 sq. m.)	<b>Secondary packaging</b> <ul style="list-style-type: none"> <li>• Two (2) controlled ambient secondary packaging production rooms, two (2) refrigerated secondary packaging production rooms, one (1) frozen secondary packaging production room</li> <li>• Cold chain labeling</li> <li>• Expiry date labeling/re-labeling</li> <li>• Just-in-time labeling</li> </ul>
Controlled ambient (15°C to 25°C)	6,921 sq. ft. (643 sq. m.)	
Scheduled drug controlled ambient (15°C to 25°C)	407 sq. ft. (38 sq. m.)	
Refrigerated (2°C to 8°C)	38 cu. ft. (1 cu. m.)	<b>Distribution</b> <ul style="list-style-type: none"> <li>• Comparator and clinical ancillary materials sourcing and management</li> <li>• Controlled/scheduled drug storage/distribution</li> <li>• Pooling of supplies</li> <li>• Reusable shipper program</li> <li>• Domestic and international transportation management</li> <li>• Import/export permit application, Importer of Record, customs clearance</li> <li>• Returns/destruction</li> </ul>
Refrigerated (2°C to 8°C)	67,825 cu. ft. (1,921 cu. m.)	
Frozen (-15°C to -25°C)	843 cu. ft. (24 cu. m.)	
Ultra-frozen (-60°C to -80°C)	52 cu. ft. (1.5 cu. m.)	

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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 for drug development, clinical trial logistics, and commercial manufacturing through our Patheon™ brand. We partner with customers in the pharmaceutical, biotech, and life sciences industries as their trusted CDMO to deliver medicine to patients faster. With more than 60 facilities around the world, we provide end-to-end pharma services across all phases of development and commercial manufacturing, including API, oral solid dose, biologics, cell therapy, mRNA, viral vectors, formulation, clinical trial solutions, logistics services, and packaging. We couple our scientific and technical excellence in these areas with a strategic partnership to provide customers of all sizes access to a global network of facilities and dedicated experts across the Americas, Europe, Asia, and Australia. Through our integrated service offerings, we provide tailored solutions to fit your unique drug development journey, accelerating your time to market.



**Discover the power of partnership and our global network.**