Summary of capabilities

Buenos Aires, Argentina

Overview:

Since 2004, we have offered specialized clinical supply chain management services for global pharmaceutical and biotech companies conducting clinical research in Argentina and Latin America. Our highly trained, multi-lingual project management team has a track record of excellence, managing 500+ clinical supply chain projects and distributing 7,000+ annual shipments to investigator sites. Close working relationships with regulatory bodies, investigator sites, contract research organizations (CROs), and local sponsor affiliates help smooth the supply chain process from end to end.

Facility facts:		Capabilities:
Opened:	2019 (original facility opened in 2004)	• Importation and exportation services including tax and duty payments and customs clearance • GMP storage at ambient, controlled ambient (15°C to 25°C), refrigerated (2 to 8°C), and frozen (-15°C to -80°C) temperatures, as well as ambient-controlled drug storage • Secondary packaging at ambient or refrigerated temperatures, as well as light-sensitive secondary packaging • In-house labeling, cold chain, and expiry date labeling/relabeling • Comparator sourcing • Clinical ancillary management • Pick and pack and distribution services that include temperature management and monitoring for local, regional, and international shipments • Clinical supply returns/storage/destruction service
Audited by:	Ministry of Health — ANMAT	
Contact info:	Av. Del Campo 1550/60 – (C1427APP) Ciudad Autónoma de Buenos Aires Argentina	
	Tel: +54 (11) 5235-9400	

Buenos Aires clinical storage, distribution, and packaging capabilities:

Storage capacity		Capabilities/services
Ambient	1,916 sq. ft. (178 m2)	**Secondary packaging** • Three (3) secondary packaging production rooms • Packaging of light-sensitive materials • Packaging in refrigerated environments • Cold chain labeling • Expiry date labeling/relabeling **Distribution** • Pooling of supplies • Comparator sourcing • Clinical ancillary materials sourcing and management • Import/export permit application, importer of record, customs clearance, domestic and international transportation management
Controlled Ambient	15°C to 25°C) 13,595 sq. ft. (1,263 m2)	
Refrigerated	(2°C to 8°C) 175,337 cu ft (4,965 m3)	
Frozen (-20°C)	10,594 cu ft (300m3)	
Frozen (-80°C)	4 units (4,000 L)	
Controlled drug	CRT Ambient) 721 sq. ft. (67 m2)	

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

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.

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