

Summary of capabilities

São Paulo, Brazil

Overview:

Our Brazil 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 Brazil and in Latin America. The highly trained project management team has a track record of excellence in the proficient management of clinical supply chain projects. Our multi-lingual team is proficient in helping global pharmaceutical and biotech companies distribute to investigator sites where the majority of shipments require temperature management and temperature monitoring.

Facility facts		Specialized capabilities
Opened	1999	 Importation/exportation services including tax and duty payments and customs clearance GMP storage at ambient, controlled ambient (15°C to 25°C), refrigerated (2°C to
Audited by:	ANVISA, CRF, COVISA	8°C), frozen (-20°C to -80°C), and cryogenic (< -150°C) temperatures, as well as ambient controlled drug storage
		 Secondary packaging in environments, including refrigerated (2°C to 8°C) and frozen (-20°C), with support for light-sensitive materials and packaging over dry ice (for below -20°C drug products)
Contact info:	Av. Jaguare 818 Unidade 29, (05346-000) São Paulo, Brazil Tel: +55 11 3769-5959	In-house labeling, including cold chain, and expiry date labeling/re-labeling
		Full comprehensive service offering of comparator and clinical ancillaries sourcing and management
		Pick and pack and distribution services that include temperature management and monitoring for local, regional, and international shipments
		Clinical supply returns/storage/destruction services

São Paulo clinical storage and packaging capabilities:

Capacity and services			
Ambient	462 sq. ft. (43 sq. m.)	Secondary packaging Three (3) CRT secondary packaging production rooms Packaging in refrigerated and frozen environments, with packaging over dry ice for -80°C products Cold chain labeling Expiry date labeling/re-labeling Just-in-time labeling Distribution Comparator and clinical ancillary materials sourcing and management Controlled/scheduled drug storage/distribution Pooling of supplies Reusable shipper program Domestic and international transportation management Import/export permit application, Importer of Record, customs clearance Returns/destruction	
Controlled ambient (15°C to 25°C)	14,571.41 sq. ft. (1,353 sq. m.)		
Controlled substance (15°C to 25°C)	624 sq. ft. (58 sq. m.)		
Refrigerated (2°C to 8°C)	9827,75 sq. ft. (913,02 sq. m.)		
Frozen (-30°C to -20°C)	7,416 cu. ft. (210 cu. m.)		
Ultra-frozen (-80°C)	199 cu. ft. (5,636 L)		
Cryogenic (-150°C)	2 tanks (22,42 sq. 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.

