Global Network

Across APAC and beyond, expanding the support to clients of all segments Worldwide



COUNTRY DEPOTS incl. partners



SHIP-TO





>200 **EMPLOYEES**



>4,000 **PROJECTS COMPLETED**





Comprehensive Range of Services

Optimise cost through agility, leveraging partners local know-how, infrastructure & assets

Clinical Trial Supply Development







Global Product Sourcing

- Ancillaries supplies sourcing
- Comparator drug sourcing
- Lab kit building
- Secondary Re-Labelling in Asia Pacific
- Regulatory support

Storage and Distribution

- IMP Patient drug kits, Ancillaries Supplies, comparator/concomitant drugs
- Validated storage facilities
- Shipment/Site Coordination
- International courier & local delivery of IMP/Ancillaries
- Specimen Handling
- Importer Of record (IOR) and customs clearance

Returns Management

- IMP Returns & Reconciliation
- Destruction Services
- Recyclable Shipper Boxes
- Multi-use temperature loggers & Data management

Clinical Reach Suite of Services



Global Product
Sourcing
CUELLIG



Clinical Storage & Distribution



Cold Chain Management



Global Project Management



Regulatory Insights



Secondary Packaging



IIT Solutions



Direct to Patient Supplies

In-Country Capability

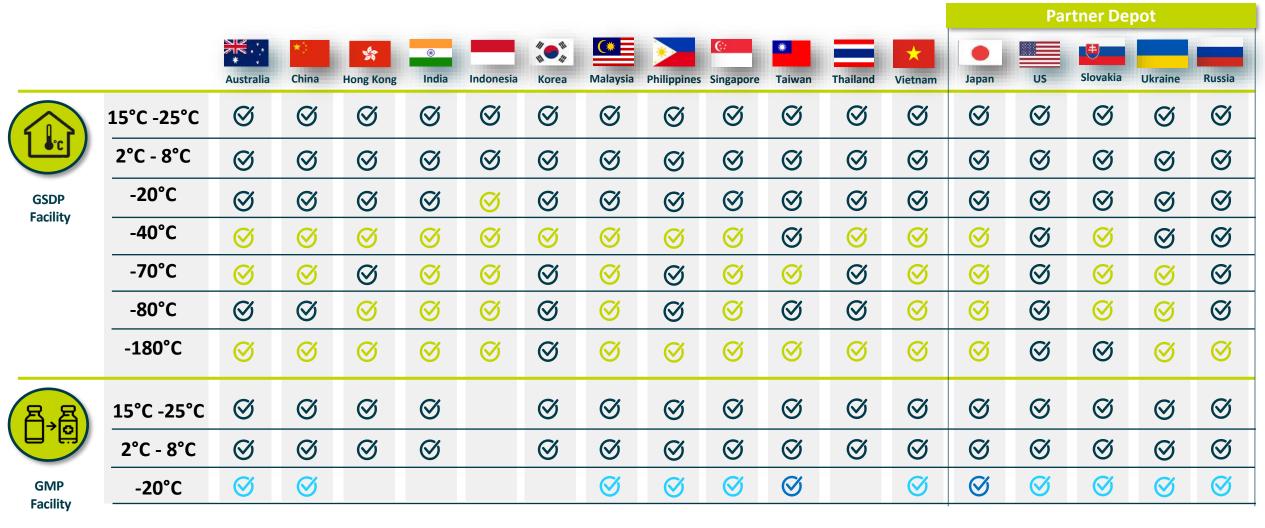
Growth as the partner of choice in APAC and beyond

		COUNTRIES	CLINICAL SOURCING	WAREHOUSE	GxP RELABELLING	CD HANDLING	DTP	IOR
ı		Australia*	V	V	V	V	V	√
		China	V	V	V			√
		Hong Kong	V	V	√		√	√
		India	V	V	√	V		
		Indonesia	V	V		V		
	APAC	Japan	V	٧	V	V		
		Malaysia	V	V	V	V	V	V
		Philippines	V	V	V	V		V
		Singapore	V	V	V	V	V	V
		South Korea	V	V	V	V		
		Taiwan	V	V	V	V		V
		Thailand	V	V	V	V	V	V
		Vietnam		V	V	V	٨	V
ROW		United States	V	V	√	V		V
	≥ ا	Slovakia	V	V	V	V		V
	RO RO	Ukraine	V	V	V	Λ		V
		Russia	V	V	√			V
		*include NZ						Δ





Extensive, Sophisticated Temperature-Controlled Infrastructure







Available Upon Request



With Time out of Refrigeration



Without Time out of Refrigeration



Evolving Needs for Comprehensive Clinical Comparator Sourcing Strategies

Zuellig Pharma recognized as the go-to supply chain partner for the clinical trial industry, and as a valued partner in solving supply chain challenges – Lead time, Speed of sourcing, Reliability

CLINICAL REACH SOURCING

US\$24 million

per annum and growing

Global Clinical Trial Supplies Market valued at **US\$2.5 billion in 2021** and is expected to grow at a decent

CAGR of 8%

PARTNERSHIP

Achieved as specialist vendor via leveraging strong lasting relationships with the

LEADING
HEALTHCARE
COMPANIES



DEPOT FOOTPRINT IN

17

APAC, EU, US

Leverages on local infrastructure and capabilities, through own depot as well local partners.



SUPPORTS OVER

5,000 CLINICAL TRIALS

(4,000 complete studies)

SMART SOURCING STRATEGIES

unrivalled access to medicine and ancillaries, working directly with global network of clinical trial sponsors and manufacturers.

We Celebrate **EcoVadis**Platinum Medal 2021-2022

We take sustainability seriously towards a **HEALTHIER**

FUTURE and are in top 1% of companies (200 industry categories, 160 countries and companies of all sizes)



Use in Clinical Trials

Setting Highest Standard

ecovadis

Quality Culture
Information Security
Integrity & Ethical
Code of Conduct
Professionalism



区部形成区

Cold Chain

Solutions

Expertise in cold-chain management with proven records. Processes & facilities aligned to highest industry standards and requirements



Dedicated Global

Procurement Professionals

responsible for the purchasing activity, posses broad understanding of the industry landscape.



With **local** and **Central** distribution capabilities, sourcing and delivering ambient, controlled temperature, refrigerated, or frozen, our footprint ensures the medicine gets to site quickly and safely.



Specialized Cold Chain Solution for Temperature Sensitive Materials

Expertise in cold-chain management with proven records



Extensive, sophisticated cold chain infrastructure and the largest*
facility in Asia (except China, Japan)



Sophisticated temperature & humidity monitoring systems track and retrieve storage data as and when required



over Facilities, Systems, Processes, Suppliers, DQ, IQ, OQ, PQ, Master Validation Plans, Validation Protocols & Reports



Highest quality cold chain infrastructure,

compliance with quality management standards (ISO9001, GMP, GSDP, GDP)



Large redressing capability in cold

chain equipped with Quality controlled environment with strict Good Manufacturing Practice (GMP) compliance



- 24/7 Temp. & Humidity Monitoring
- Alarm system for temp. excursion
- Walk in chamber
- Back up generator
- Yearly validation

cold room Processes & facilities aligned to highest industry standards & requirements







Secondary Repackaging Facility

Enhancing Quality and Increasing Capacity for Redressing



CONTROL

- Dedicated access-controlled areas
- Approved and audited SOP
- Lines are arranged to form process work center for better supervision and order allocation



STORAGE

- > Specialized and controlled packaging material storage
- Segregated area for repackaging of cytotoxic products and controlled substances



AREA*

- Authority's approved manufacturing site
- Temperature control and monitoring, humidity monitoring
- Relabeling condition: Controlled Ambient 15-25°C, Cold Room 2-8°C, Frozen -20°C



CERTIFICATION & CAPABILITIES*

- > cGMP compliant facility, re-label and repackage clinical trial supplies that are conformant to international standards, local regulations and applicable SOPs
- ➤ Relabeling, label/temper seal printing, manufacturer kit box, kitting, JIT relabeling, etc.





Direct to Patient Solutions

We offer3 type of services:



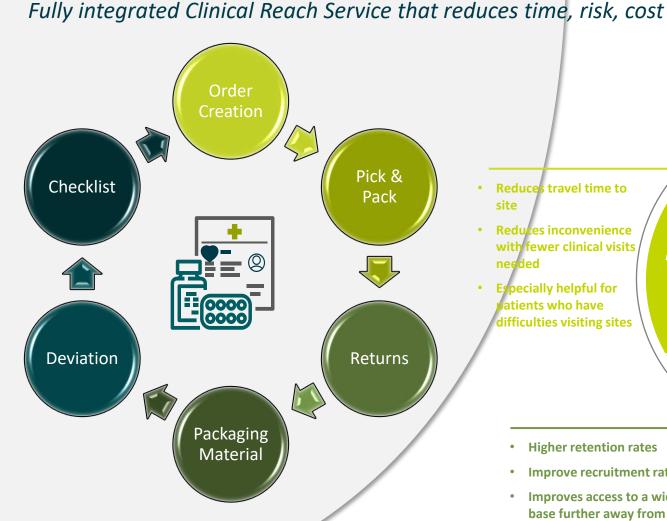




Depot to Patient

Site to Patient

Direct from Patient



- Reduces travel time to site
- Reduces inconvenience with fewer clinical visits needed
- Especially helpful for patients who have difficulties visiting sites



Benefits of Dtp

Reduces the number of visits to clinical

healthcare sites

- **Reduces facilities** management
- Retain patient for clinical trial

- Higher retention rates
- Improve recruitment rate and time
- Improves access to a wider patient base further away from sites
- Increase data quality



making healthcare more accessible