KANISHK INTERNATIONAL – GLOBAL PHARMACEUTICAL, CLINICAL RESEARCH TRIAL SUPPLIES & LOGISTICS

GLOBAL HEALTHCARE GROUP

Expertly bringing significant change & simplification to clinical trials & patient healthcare supply chain.

K-INTL's GLOBAL CLINICAL RESEARCH & PHARMA COLD SUPPLY CHAIN PROVIDES:

- Planning, forecasting and mapping all activities to optimize the Pharma SCM & ancillary supplies strategy
- 'Real-Time' Monitoring and Auto Reports of Pharma Cold Supply Chain on IoT / Cloud Networks
- Proactive project management to drive the success of your project
- Competitive sourcing of Clinical trial material, medical equipment / devices / diagnostic materials and ancillaries
- Network for global storage & distribution, inventory control, returns and destruction and final clinical trial material reconciliation
- Global trade compliance (import / export) and regulatory support for ancillary supplies
- Proven track record of work efficiencies
- Business Intelligence and Analytics to meet satisfaction of stakeholders

PARTNERING WITH CUSTOMERS TO IMPROVE PATIENT HEALTHCARE OUTCOMES

We are committed to serving our customers' evolving drug development & medical treatment needs. A single trusted source – with the experience, resources, and capabilities that cover the full spectrum of services across the supply chain, from pre-formulation manufacturing to global logistics and distribution, **K-Intl GCSC** exhibits transformation of the global clinical trial lifecycle.

YOUR CLINICAL SUPPLY: MANUFACTURE / ANCILLARY STRATEGY STARTS AT PLANNING AND FEASIBILITY

K-Intl has a strong local presence and is known for its professional and efficient conduct. Companies undertaking bioequivalence / bioavailability (BE/BA) studies phase I-IV and medical device clinical trials benefit from our expertise in site selection, study feasibility, and monitoring services, as well as our knowledge of regulatory affairs, EXIM, warehousing, and Pharma Cold Supply Chain. We provide supplies of drugs and ancillaries for a worldwide study that requires in-depth knowledge of the export & import regulations in each country regarding those supplies around the globe. It importantly requires global awareness of manufacturer's products and materials availability. Meeting study milestones and timelines is a high priority when formulating a strategy for clinical supplies. **K-Intl** offers global expertise to plan your drug & ancillary supply sourcing and distribution strategy the moment you commence the program / study. We encourage sponsors to engage **K-Intl** at protocol development or earlier to plan timelines and mitigate risks. We exhibit our proactive approach to ensure strategic planning and execution is integrated into the entire trial process. **K-Intl** can organise drug manufacturing, purchase supplies on a global or local basisas required for each study.

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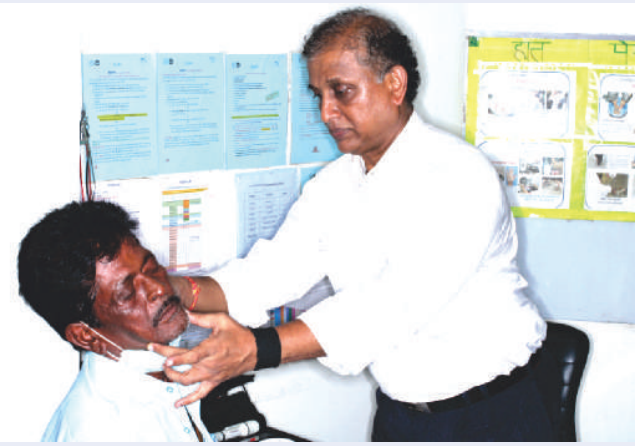
'Nurturing Excellence in Pharma - Clinical Supply Chain & Cold Chain Management'

SIMPLIFIES END-TO-END ANCILLARY SUPPLIES DELIVERY BY PROVIDING:

- Proactive clinical logistics leadership
- Central material flow management, leading to significant shipping cost savings and an optimized supply chain
- Centralised monitoring for seamless SCM functioning & maintain OTIF levels
- Proven time and process efficiencies
- Inventory transparency
- Business intelligence



Dr. V V Pai - Dermatologist & Leprologist

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For program level strategies with projects at multi centric locations / countries we have participated to design a pooled supply inventory strategy that drives efficiencies and cost controls. This enables our valued customer to balance the cost of ancillary supplies, delivery logistics and storage, to ensure that the project runs lean while guaranteeing treatment and supply continuity.

Whether the customer is a large, global pharmaceutical company with decades of experience, or an emerging biotechnology company with no in-house supply chain expertise, **K-Intl** 's GCSC experts can help you plan and execute your entire clinical trials supplies process to make it more efficient and cost-effective.

PROCUREMENT, CONTRACTUAL MANUFACTURING AND PACKAGING STRATEGICALLY

K-Intl enables the Customer to determine the right source of procurement of drugs, customized contractual manufacturing, ancillary supplies; totally depending on the location of trial sites, and the patients being recruited. Through joint working we help customers to determine your ancillary supply strategy and vetting the benefits of sourcing locally or globally for your trial. We provide both local and global sourcing and procurement support and can manage suppliers, contracts, rate cards, scorecards and evaluate performance.

K-Intl GCSC amalgamates supply of drugs, ancillary and lab supplies; by centralizing manufacturing, sourcing, and shipments; we reduce the risk of loss of goods, thereby maintaining vision of control on shipping costs and increasing inventory transparency.

Focused on Quality, Innovation and Manufacturability – we cater to a range of packaging of solid dosage pharmaceutical products, with anti-counterfeiting features to safeguard the brands and consumers.

SOLVING INDUSTRY DISTRIBUTION AND STORAGE CHALLENGES AROUND THE WORLD

K-INTL has international associations of logistic partners at multi locations with depots designed specifically for Pharma SCM and operates facilities to store and distribute Pharma & clinical trial materials. We have expertise to maintain an uninterrupted supply chain throughout Americas, Europe, Africa and the Asia Pacific Region.

We enable the import - export of pharma, medical equipment, prototype systems and devices; with expertise to manage Customer projects in seamless form even at remotely located sites.

In addition, **K-INTL 'Real-Time' Monitoring Solutions** of: Pharma Cold Chain / ancillary supplies, *e*-Logistics to manage all inventory's, stocks and expiration dates.

K-INTL SMO – SITE MANAGEMENT ORGANISATION FOR CLINICAL TRIALS

K-INTL SMO provides site management organization (SMO) solutions for clinical trials of treatments. With outsourcing, insourcing, and procurement abilities, the company's GCSC team specializes in supporting clinical operations, project management, and quality assurance activities under directions from the principal coordinator / investigator (PC / PI).

As a part of its procurement skills, **K-INTL SMO** sources cost-effective comparator drugs and standard-of-care medications, as well as ancillary supplies and equipment such as freezers, 'Real-Time' monitoring devices, and passive shipping solutions.

K-INTL association with strategic partners at regional depots, along with our intelligent, fully customizable ***e***-Logistics tools provide full clinical supply chain transparency, inventory management, and ancillary supply traceability for end-to-end sample management across the entire life cycle of your trial. Services are carried out with full accountability and maximum flexibility.

KANISHK INTERNATIONAL - CLINICAL TRIAL SUPPLIES & LOGISTICS: ANCILLARY SUPPLIES: STRATEGICALLY, GLOBALLY INTEGRATED

K-INTL hasa proven track record of a trusted partner for the complex development journey required of bio-pharmaceutical and medical device companies.

For the past 10 years, **K-INTL** Clinical Trial Supplies & Integrated Logistics has provided high-quality, centralized coordination of clinical trial supplies, pharma / laboratory logistics services and ancillary supplies. Available individually or in any combination, **K-INTL** assures that medications / supplies reach your sites and your patients on time.

Dr. Anju Wakade - Research Scientist (Microbiology)

DEPOT ASSET MANAGEMENT – Clinical trial storage services

K-INTL is in process of setting up clinical trial specific warehouse under accreditation from CDSCO / DCGI, India – to provide stand-alone services for clinical trial drugs, medical diagnostics & devices, while assisting in making import licence applications to CDSCO, customs clearance for importing and exporting the items and distribution (via **K-INTL GCSC)**.

We are associated with pharma storage warehouses with huge capacity and maintain 'Real-Time' monitoring of controlled temperatures of +15°C-25°C, +2°C-8°C, -20°C, and -80°C. The state-of-the-art facilities are in accordance with GDP, CFR-21 (Part 11), and FDA.

CLINICAL SUPPLY CHAIN BUSINESS ANALYTICAL INSIGHTS: ONLY BY **K-INTL**

Health Group of Clinical Trial Supplies & Logistics analytics gives you the best visibility and control over your pharma, devices, & ancillary supply logistics. Our state-of-the-art ***e*** - Logistics solutions are the key to trackall ancillary supplies from vendor and sponsor to depot and clinical site through return and destruction during the course of your study and provide insight into operational supply chain performance.

Combined with **K-Intl** expertise, it enables companies to build supply chain intelligence that can predict issues, mitigate risks, and implement actions to optimize OTIF levels, reduce costs and enhance future trials.

ABOUT **K-INTL** HEALTHCARE GROUP

Since incorporation, **K-INTL** has delivered key CRO services to pharmaceutical / medical diagnostics – devices sectors. The company's innovative, globally oriented dedication benefits both patients and clients. The global outreach and associations help to manage all trial management processes in early and late - phase studies.

K-Intl Healthcare Group provides development strategies, assisting medical device development from discovery to invention processes and clinical trials. The company's high-value services provide the flexibility, individualisation, and easy-access offerings required for collaboration with local and multinational pharmaceutical companies, health centers – hospitals, doctor associations, and health funds.

To learn how KANISHK's Ancillary Supplies Experts can help you to optimise clinical trial supply chain,

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