CRITICAL SUPPLY CHAIN STAGES

PLANNING

Forecasting and strategy

SOURCING Drugs, ancillary

supplies, and lab

materials

MANUFACTURING Packaging and lab kit production

STORAGE

Warehousing and

inventory control

Managing shipments, imports,

DISTRIBUTION

and exports

SITE Storage and inventory control, sample management, and

data processing

RETURN Managing shipments, final clinical trial

materials

reconcilliation

DESTRUCTION Recycling / destruction

ANALYTICS Data processing and analysis

INSUFFICIENT COMMUNICATION

Poor coordination of the clinical trial materials supply chain can result in both duplicated effort and action gaps. Errors and changes in strategy mid-trial require frequent plan revisions.

PROCESS DELAYS

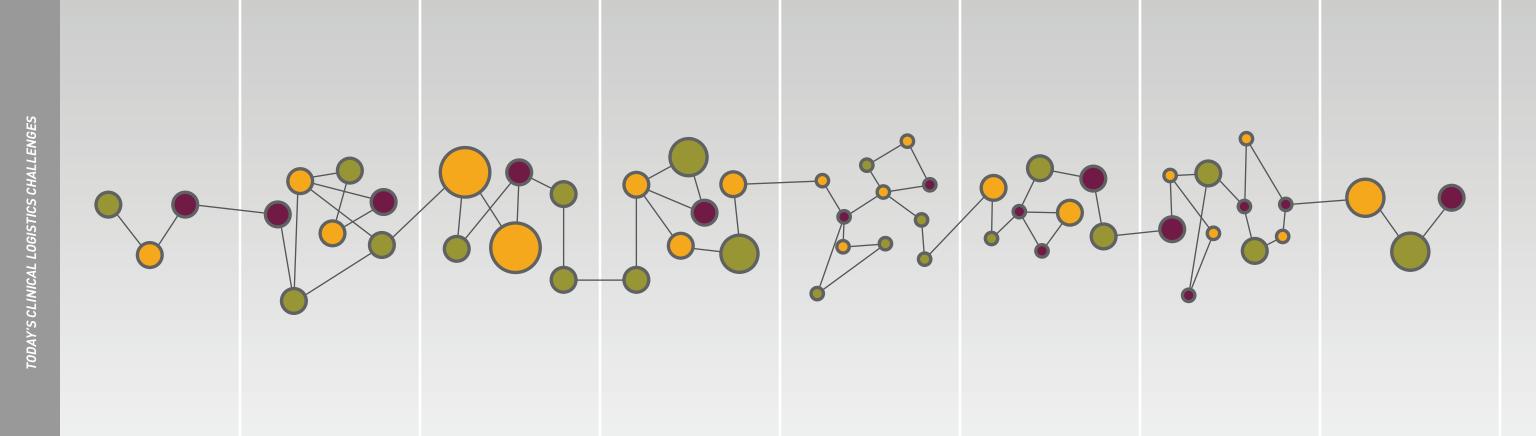
Since materials suppliers are relatively disconnected from one another, the overarching implications of a supply chain issue may not be apparent or clear, often resulting in delays at trial sites.

INFORMATION GAPS

Very often, overall study timelines are endangered by blind spots inherent in today's common practices, leading to out-of-stock situations that cause patient recruitment delays, and unexpected expiration of products that result in patient safety issues.

HIGH RISK OF COST OVERRUNS

Inefficiencies in the system, the difficulty of identifying and addressing all cost drivers, and the potential expenses of corrective actions frequently drive costs well over anticipated budgets.





No comprehensive clinical supply chain intelligence. New trials must start without the benefit of accumulated knowledge and experience.

CLINICAL TRIAL SUPPLIES

ANCILLARY SUPPLIES

AST

CLINICAL LOGISTICS LEADERSHIP

PAREXEL provides a single point of contact for clinical trial materials management. This efficient approach helps sponsors.

ACCELERATE SITE READINESS

Intelligent forecasting and developing the best logistics strategy, combined with local expertise for importing clinical trial materials, are key success factors to getting sites online swiftly.

MINIMIZE SUPPLY CHAIN ISSUES

Centralized coordination of the entire supply chain reduces the risk of issues significantly, and allows for rapid response when they arise.

REDUCE PATIENT SAFETY RISKS

By overseeing all process-critical materials supply tasks, many safety risks such as out-of-stock or unforeseen product expirations are mitigated.

CREATE BUSINESS INTELLIGENCE

A centralized eLogistics Service, including IVRS/IWRS and simulation tools, creates a significant knowledge base of historical data for planning future

OPTIMIZE OVERALL COSTS

PAREXEL's unique approach to planning, execution management, and supplier monitoring provides significant cost savings along the supply chain, gets products to patients faster, identifies overspending budget risks upfront, and monitors each budget category centrally.



CLINICAL **SUPPLY CHAIN ANALYTICAL** INSIGHTS, ONLY BY PAREXEL



Supply chain intelligence s developed through the unique combination of PAREXEL expertise and our state-of-the-art eClinical and eLogistics solutions. We use this intelligence the process, to anticipate issues, develop solutions optimize supply levels, reduce costs to clients, and enhance future trials by tightly integrating with the entire trial process.

PAREXEL® YOUR JOURNEY. OUR MISSION.™

CLINICAL RESEARCH SERVICES: CLINICAL LOGISTICS

SIMPLIFYING END-TO-END GLOBAL CLINICAL LOGISTICS

www.PAREXEL.com/clinicallogistics

Clinical trial globalization creates significant logistical challenges. Effectively managing the clinical trial materials supply chain requires strategic oversight and up-to-the-minute tactical knowledge—plus a significant commitment of resources—to maintain reliable deliveries and avoid costly delays. Many organizations would benefit from a tight integration of supply chain information yet few are equipped to manage the increasing complexity of the trial materials supply chain.

PAREXEL can turn clinical trial complexity into clinical logistics intelligence. As your dedicated partner, PAREXEL ensures that logistical planning and execution is seamlessly integrated throughout the entire clinical trial life cycle to minimize disruptions and delays.

With strong support from PAREXEL, our clients and sites spend less time worrying about supplies and logistics, and are able to focus their attention on critical trial issues.