EUROMED PHARMA

SERVICES TO CLINICAL TRIALS

Euromed Pharma provides strategic and integrated solutions as one single provider of services to healthcare in clinical trials with one single point of contact and global approach.

1 COMPARATOR SOURCING

Euromed Pharma Comparator sourcing strategic team committs to support Clinical trials by providing :

- Comparators
- Rescue medications
- Ancillary materials
- Concomitant and background medications

with:

- certificate of Analyses (CoA),
- certificate of Conformity (CoC)
- TSE/BSE statements,
- batch release certificates (BRC),
- certificate of origin (CoO)

As an **Importer of record (IOR)**, we act as a bridge between the customs authorities and our customers, ensuring the smooth and efficient access of products into the destination country, handling the whole documentations.

OUR COMPETITIVE ADVANTAGES

Clinical Supply Chain Consulting

Preliminary Assessments to successfully plan your trials:

- Strategic planning – market accessibility intelligence
- Clinical Supply management forecasting tools

Operating Globally

- Tailor-made solutions and customized service (SPOC).
- Global procurement: access to 80K+ SKUs
- Sourcing comparator drugs in 60+ countries via our global wholesale network for cost savings and access to key drugs.

Keeping the supply chain short

- Strategic and integrated solutions.
- Flexibility, efficiency, and supply continuity throughout your trial.
- Strong partnerships with generic and biosimilar manufacturers for significant cost savings.

2 IMP MANAGEMENT

At Euromed, we provide end-to-end IMP management for clinical and commercial needs. Through an integrated approach, including manufacturing, storage, distribution and temperature protection, we support our customers to face IMP challanges, reducing costs and improving efficiency.

QUALITY

- Good Manufacturing Practices (GMP)
- Quality Control (QC), Quality Assurance (QA), Qualified Person (QP)

+

EXPERTISE

Long-standing and trusted relationships with well-known pharmaceutical manufacturers, IRCCS and hospitals.

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Development of **innovative ideas** for clinical trial from **phase I to phase IV**. Our approach is in **teamwork** with project managers and clinical experts to **identify and optimize the management and implementation of IMPs for clinical trials.**

IMP MANUFACTURING

Specialized in placebo and drug capsules & Innovative IMP labelling techniques

IMPs Manufacturing

- Blinding of IMP: both semi-automatic and manual capsules preparation
- Capsules Manufacturing
- Batch release & certification by QP

IMPs Labelling

- Label design
- Multilingual labeling
- Re-labeling in case of changes
- Compliance with national drug laws
- Tear-off label for subject compliance and drug accountability

IMPs Packaging

- Primary packaging for solid oral formulations
- Secondary packaging
- User-friendly materials
- Tailored packaging and labeling

IMPs Services

- Emergency Envelops, international Master Label Forms and IMPD
- Importation of IMP from extra-EU countries and production of QP declaration
- Narcotics management for CT

3 STORAGE & SHIPMENT

Competence, high responsiveness, flexibility and a wide range of options for Clinical trial storage and distribution in Italy, in Europe and all around the world.

STORAGE

- Manufacturing and packaging: +15°C/+25°C, +2°C/+8°C, and -15°C/-25°C to **-80°C**.
- Analytical laboratory with temperature-controlled conditions.
- Temperature and humidity-controlled primary packaging areas.
- Segregated storage area for narcotics, with 24/7 monitoring.

SHIPMENT

- Shipment prepared within 24 hours, delivered in 24-72 hours across Europe, with temperature-controlled
- Just-in-time delivery (Italy: 12-24h; Abroad: 24-72h)
- Validated boxes for temperature-controlled shipments and specific Clinical Trial forms.
- IMP resupply managed via IXRS

3 RETURN & DESTRUCTION

Sponsor support service for the management of the final part of the clinical trial life cycle. Management and reporting of withdrawal service for Clinical Trial expired and not expired drug.

RETURN

- Returns of expired and unexpired drugs managed by a qualified external provider
- Option for storage and management of returns by the customer
- Reconciliation and control of returned materials as per customer specifications
- Storage in a segregated area awaiting destruction or removal by the customer

DESTRUCTION

- Transfer to the site authorized for the drug destruction
- Preparation of specific study form
- Certificate of destruction issued

GET IN TOUCH

EUROMED PHARMA: 11 WORLDWIDE LOCATIONS

EUROMED PHARMA
TOGETHER THROUGHOUT YOUR DRUG'S LIFECYCLE

Ireland Dublin
Italy Naples, Milan
Germany Berlin
France Paris, Lyon
Spain Barcelona
Portugal Lisbon
UK London

EUROMED PHARMA US
a Petrone Group company
North Carolina Raleigh

EUROMED PHARMA APAC
a Petrone Group company
Singapore

EUROMED PHARMA PORTUGAL
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