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CLINICAL TRIAL SUPPLY

PACKAGING AND LOGISTICS

EXPERTISE

- 30 years of experience in Clinical Trial Supply
- Worldwide experience
- Cold chain capacities
- QP release
- Double-Blind management
- Controlled drugs services

Drug Product

Medical Device

SERVICES

Primary Packaging

- Placebo development
- Blisters of sterile (vials, syringes) and non-sterile forms (capsules, tablets)
- Bottles, sachets...
- Over encapsulation

Secondary Packaging - Labeling

- Customized labels / Leaflets design and printing
- Boxes and wallets / Patient kit design
- Randomization list and decoding envelopes
- Ancillaries supply and kits
- Comparator supply, blinding and labeling
- Extension of expiry date

QP Services

- Import, testing and certification and/or batch release
- Site audits to support the QP declaration / QP agreement
- GMP certification | Final batch release

GMP Storage (APIs, DP, IP)

- 15 - 25 °C | 2 - 8 °C | - 20 °C | - 80 °C

Distribution

- "Just in time" labeling
- Worldwide shipments to clinical sites and depots
- Returns and destruction

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CLINICAL TRIAL SUPPLY

PACKAGING AND LOGISTICS

FACILITIES

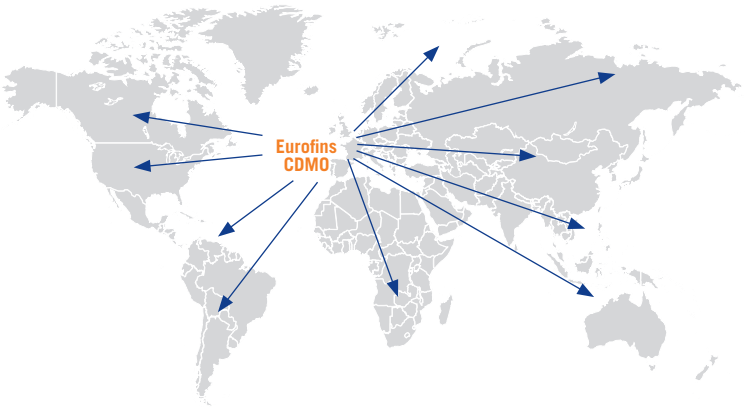
- 2 dedicated sites for CTS activities
- 100+ people
- 5 ISO 7 suites / 3 ISO 8 suites
- 10 secondary packaging suites at RT
- 3 secondary packaging suites at 2/8°C
- Storage capacities:

T°	+15 / +25 °C	+2 / +8 °C	-20 °C	-80 °C
m^3	560 m^3	146 m^3	2.6 m^3	0.5 m^3
ft^3	19 775 ft^3	5 156 ft^3	92 ft^3	17 ft^3

- Controlled drug areas

ANSES ANSM GMP GDP Crédit Impôt Recherche

DISTRIBUTION: GLOBAL CLINICAL SUPPLY SOLUTIONS



- Certified depots in US
- Access to worldwide depots through our courier partners (Asia, Russia, Latin America...)
- Global clinical supplies management for US and EU companies at controlled temperature

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