

CLINICAL TRIAL SUPPLY

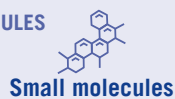
Packaging and logistics

EXPERTISE



- 30 years of experience in Clinical Trial Supply
- Worldwide experience
- Cold chain capabilities
- QP release
- Double-blind management
- Controlled drugs services
- Patient kit design
- Direct-to-Patient logistics

MOLECULES



PRODUCTS



6800
shipments
per year

530
batches
released
per year

SERVICES

Secondary Packaging - Labeling

- Customized labels / Leaflets design and printing
- Customized kitting solutions (boxes, wallets)
- Randomization list and decoding envelopes
- Ancillary supplies
- Comparator supply, blinding and labeling
- Extension of expiry date

QP Services

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

Additional Services : Primary Packaging

- Sterile / non-sterile development and manufacturing (Drug Product / Placebo)
- Blistering
- Over encapsulation

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

CLINICAL LOGISTIC SERVICES



Hospitals / clinics /
clinical trial centers



Direct-to-Patient



CLINICAL TRIAL SUPPLY

Packaging and logistics



- Clinical packaging and logistics
- Sterile manufacturing
- Non-sterile manufacturing

FACILITIES

- 1 dedicated site for CTS activities
- 100+ people
- 8 secondary packaging suites at RT
- 1 secondary packaging suite at +2°/+8°C
- Storage capacities:

T°	+15°/+25°C	+2°/+8°C	-20°C	-80°C
m ³	800 m ³	150 m ³	5 m ³	2 m ³
ft ³	28 250 ft ³	5300 ft ³	177 ft ³	71 ft ³

SOON 2024: Additional storage capacities available

- Controlled drug areas

ANSES ANSM **GMP** **GDP**



DISTRIBUTION: GLOBAL CLINICAL SUPPLY SOLUTIONS

- Global management of your clinical supplies
- Access to qualified worldwide depots
- Continuous sourcing of new logistics partners
- Access to worldwide depots (Asia, Australia, Latin America...)



CTS01



Full-services CRO

from study design to Clinical Study Report



Why working with us ?

- Private phase I unit: 60 beds, ambulatory area
- More than 30 years of experience
- Drug, food and medical devices
- In-house multidisciplinary experts
Pharmacologist | Pharmacist | Physicians | Nurses | Project manager | ...
- 30 patient studies over the past 5 years in 70 investigator sites
- Human scale CRO with Eurofins support
- GCP certified
- Flexibility, rapid decision making and agility



A BROAD EXPERIENCE IN VARIOUS THERAPEUTICS FIELDS (%)

