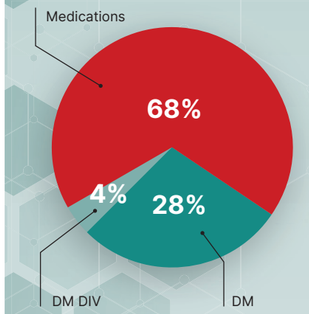
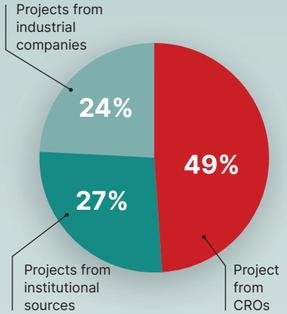
Médi-link since 2016

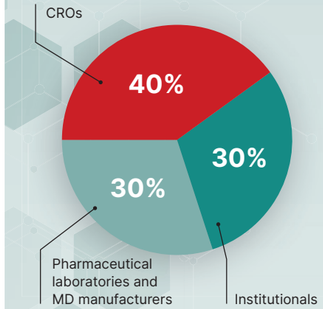
STUDY BY PRODUCT TYPE



CLIENTELE STUDY- TYPE



CLIENTELE TYPOLOGY



THERAPEUTIC AREAS

- Oncology
- Infectiology
- Ophthalmology
- Orthopedics
- Hepatitis
- Onco-Hematology
- Gynecology & Obstetrics
- Hematology
- Inflammatory Diseases
- Respiratory/ENT
- Venous Compression
- Gene Therapy
- Pediatrics and Neonatology
- Urology

Our goal by 2030 is to become a key leader in providing support for hospitals, pharmaceutical laboratories, and medical device manufacturers.

Medi Link
Transformons vos essais

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Guarantee the quality of your clinical studies and ensure the success of your initiatives

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Medi-Link is a human-scale CRO that has been in the healthcare sector for over 10 years. We act as experts for investigator sites to secure and optimize your clinical projects, complementing your teams or the teams at investigator sites.

A true «HUB» bringing together qualified and recognized partners, we are able to offer you services that go beyond those of a classic «Full-Service CRO» through flexible, agile, and expert structures in their respective fields. This allows us to tailor the structure and budget of our services precisely to your needs without multiplying your points of contact.

OUR MISSION

To restore trust among the various stakeholders, ensure the quality of your clinical studies and guarantee the success of your initiatives, even under critical time constraints.

OUR VALUES

Reliability in supporting biomedical research for the well-being of patients, while fostering a work environment that respects the balance between professional and personal life.

1 Reliability

Expertise, Quality of Work, Guidance, Client Satisfaction, Physician Satisfaction, and most importantly, Patient Satisfaction.

2 Reactivity

Clarity of missions, Regional employees, Availability, Local networks, Quick responses, Well-defined roles

3 Commitment

Team spirit, Involvement, Adherence to indicators, Mutual support, Kindness, Quality of work life, Work life balance

Services

1 RESEARCH DESIGN

Methodological Expertise: Research typology, calculation of required sample Size, protocol writing

Design and Writing of Documents: Protocol, investigator brochure, Informed consent, data collection form and case report form

Configuration of Electronic Data Collection Tools: eCRF, ePRO, eConsent.

2 REGULATORY SUBMISSION

ANSM, CPP, CNIL, Health Data Hub

ClinicalTrials.gov CTIS

ARS, CNOM (IDAHE2 Plateform)

3 PRIVILEGED SELECTION OF INVESTIGATOR NETWORKS

Qualification of Research Sites : Hospitals, private clinics, private practices, pharmacies.

Feasibility on the ground : Provision of the most suitable centers for your project (therapeutic area, scientific reputation, recruitment potential, human resources, and equipment).

Financial agreement: Negotiation of hospital costs/overheads and drafting of financial agreements.

4 CONDUCT AND MANAGEMENT OF CLINICAL TRIALS

Project Management: Coordination and oversight of the trial, recruitment management and optimization, trial safety monitoring.

Site Management: Selection and qualification of sites, site initiation visits, monitoring visits (on-site and remote), monitoring reports, close-out visits, investigator support.

5 DATA MANAGEMENT AND SCIENTIFIC VALORIZATION

Data-Management & Statistics: Plan de data-management, Plan d'analyses statistiques, traitement des données, analyses statistiques, rapport statistique.

Medical writing: Study report, publication, poster, results valorization, communication document writing.