

Pharm Research :

- **20 years experience in the clinical trial field**
- Experimented and graduated staff
- Management of Clinical trials
- Partnership with national and international pharmaceutical laboratories, Hospital and Clinic
- **Monitoring and audit**

Services provided ?

Pharm Research offers a full range of services :

- Study Coordinators/CRA on site
- **Management of phase I to phase IV clinical trials** : from site initiation visit to study closure.
- **Management of interventional and observational clinical trials**
 - Accurate feasibility and Initiation site visit
 - Support to investigators: we provide staff to facilitate the recruitment of patients by handling the administrative and management of trials
 - **Simplification** of trials supply and documentation
 - Logistical
 - CRF/eCRF completion with **Quality data** from experienced GCP-trained staff, leading to medical data quality
 - Resolution of queries and deadlines respect
 - Management and follow-up of adverse event, serious adverse event, and end point
 - Transmission of pharmacovigilance, materiovigilance, reactivovigilance data
 - Monitoring visit and report
 - Patient recruitment :

For each trial, recruitment strategies are studied by our staff and adapted to the case (Worksheet, Standard Operating Procedures, Logistic management).

To give the study the best chances of success, Pharm research develops a team spirit involving the principal investigator, the sub-investigators, the study nurse, the CTA and the patients.

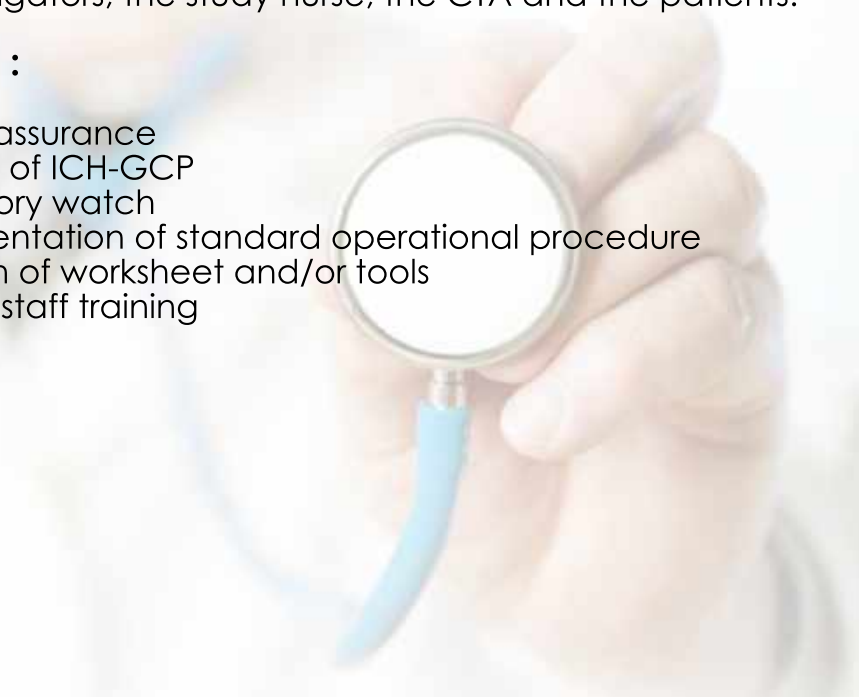
Expertise :

- Quality assurance
- Respect of ICH-GCP
- Regulatory watch
- Implementation of standard operational procedure
- Creation of worksheet and/or tools
- Regular staff training

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- Our employees have a solid experience in clinical trial (Clinical Research Engineering, Master's degree in Clinical Trials, Master 's degree of Clinical Pharmacy, Master's degree of Quality Assurance and Audit).
- We are focused on Phase I-IV clinical trials both in promotion and monitoring.

Gestion administrative :

- Contracts
- Budget negotiation
- Respect of regulatory standards

Therapeutics areas :

Cardiology
Medical devices
Oncology
Nephrology
Hematology
Nutrition
Geriatric
Endocrinology : Diabetes type II, hyperlipemia
Psychiatry
Gastro-enterology

Site Management :

- Deadline and timeline
- Dedicated trained and experienced staff for each site.
- Audit preparation
- Other procedures and services may be available according to practical needs.

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