Leading Clinical Research

We will take you through the labyrinth of clinical trials directly to your target – innovative, safe and effective drug or medical device

Company presentation

2018

LCR

Contract Research Organization
with HQ in Prague, Czech Republic

Operations in Europe
with focus on Central and Eastern Europe

Our core region (CEE) is known for:

- ✓ Qualified, highly motivated investigators
- ✓ Excellent recruitment
- ✓ High data quality
- ✓ Cost-effectiveness
- ✓ Trial-friendly regulatory climate

Services

Services provided - customized to clients´ needs

- ✓ Study design / protocol development
- ✓ Regulatory submission
- ✓ Subject recruitment
- ✓ Clinical conduct
- ✓ Medical writing
- ✓ Data management
- ✓ Statistical analysis
- ✓ Study report
- ✓ CRA and/or Investigator GCP and ISO 14155 Trainings

LCR Staff

- ✓ Project Managers
- ✓ **CRAs - all with University degree, regularly trained in GCP and ISO 14155**
- ✓ Medical Writers
- ✓ QA/QC Managers
- ✓ Data manager / Statistician
- ✓ Training division
- ✓ External group of experts covering all therapeutical areas

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Therapeutic Area Experience:

- ✓ Cardiovascular Diseases
- ✓ Gynecology/Urology
- ✓ Ophthalmology
- ✓ Rheumatology
- ✓ Oncology
- ✓ Dermatology
- ✓ Pediatric and Vaccination studies
- ✓ CNS incl. Neurosurgery
- ✓ Respiratory Diseases
- ✓ Gastroenterology, Hepatology
- ✓ Diabetology
- ✓ Orphan Drugs
- ✓ Infection Disease

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Quality standards

Quality management system certified according to ISO 9001:2008 for:

- ✓ Clinical Trials of Medicinal Products
- ✓ Clinical Investigations of Medical Devices
- ✓ Non-interventional Trials
- ✓ Medical Writing
- ✓ Trainings

- **Functional QA/QC system**, relevant SOPs in place
- **Regulatory Authority Inspections** on Clinical Trials and Clinical Investigations repeatedly without any findings

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We reach beyond Europe

AICROS ASSOCIATION OF INTERNATIONAL CONTRACT RESEARCH ORGANIZATIONS

- ✓ **AICROS is a network of local & well established CROs businesses providing full range clinical research services on a global scale.**
- ✓ Region covered so far includes almost **all European countries, Israel, North and South America and Japan**
- ✓ Four major principles:
  - Integrated training curriculum
  - Bi-annual members‘ audits to secure the high level of quality in services provided
  - Anticorruption procedures
  - Sharing financial results to show financial stability

LCR clients

Pharmaceutical companies

Medical Devices Manufacturers

Bio-tech

CROs

Since 2009: 67 projects, more than 600 sites and 6.000 subjects

WHY LCR?

- ✓ Deep knowledge of local investigators and clinical trials I-IV
- ✓ Very good reputation with regulatory authority, 100% approved trials
- ✓ Very stable teams: low CRA turnover, management team stable since the company foundation
- ✓ Experience with combination products (that combine drugs, devices, and/or biological products)
- ✓ Highly customized projects and individualised budget calculations
- ✓ 100% repeated business
- ✓ Proven recruitment record

Last but not least

"Details make the difference between good and excellent clinical trials. We pay close attention to details to achieve the highest level of excellence.“

Jiri Paseka, MD

Managing Director and Founder

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