

Central and Eastern Europe and the Middle East



Setting Standards for Clinical Trials

Established in 1996, Quintiles was the first global CRO in Central and Eastern Europe and the Middle East. And today, with approximately 1,500 employees in offices throughout 18 countries, we give you access to the right patient groups at the right time via a regional population of almost 580 million.

Our multiple sites make reaching patients faster and easier, even beyond our established offices. Quintiles' strong, long-term relationships with more than 14,000 investigators facilitate recruitment. Our successful collaboration with leading sites and experience with competent local authorities speed up your trials.

Enlist Quintiles as your ally in *the New Health* — the rapidly evolving world of biopharma. We have the proven resources and expertise to help you seize emerging opportunities and deliver on your objectives.

State-of-the-Art Technologies

- > Medical facilities that are among the world's best offer you high quality and enhanced efficiency
- > Web-based and mobile communication tools facilitate clinical monitoring and follow-up
- > Adherence to Quintiles' high-quality global operating procedures and FDA/EMA (formerly EMEA) standards

Prevalent Diseases

- > Cardiovascular disorders
- > Neurologic and psychiatric disorders
- > Cancers
- > Respiratory / Pulmonary disorders
- > Diabetes

Operational and Delivery Excellence

- > The proven market leader in Central and Eastern Europe and the Middle East, Quintiles has the depth of resources and expertise to successfully deliver your trials
- > We have delivered more than 447 studies in the last five years, involving over 144,000 patients
- > We manage your studies efficiently and effectively due to:
 - Our in-depth knowledge of clinical trial requirements and trial logistics in each country and locale
 - Our highly educated, English-speaking clinical specialists (CRAs, CTLs, PMs, investigators); more than 70% have master's degrees or Ph.D.s
 - Thorough training programs and a retention rate of over 90%
 - Deep understanding of different cultures, regulatory requirements and healthcare environments
 - Access to and relationship with network of clinical research centers of excellence
 - Dedicated Access to Patients (ATP) program

Quality Data

- > Data capture and management throughout the region complies with ICH-GCP, EU Clinical Directives and FDA regulations
- > EDC management services adhere to global standards

Key Facts

Regional

- > A large population spread across 30 countries, including:
 - Central and Eastern Europe – Austria, Bosnia & Herzegovina, Bulgaria, Croatia, Czech Republic, Estonia, Greece, Hungary, Latvia, Lithuania, Macedonia, Malta, Montenegro, Poland, Romania, Russia, Serbia, Slovakia, Slovenia, Ukraine
 - Middle East – Egypt, Israel, Jordan, Kuwait, Lebanon, Oman, Qatar, Saudi Arabia, Turkey, United Arab Emirates
- > Established economies in many countries and growing economies in others
- > Centralized healthcare systems with highly motivated investigators and clinical research personnel
- > An estimated \$25 billion pharma market with a 20% annual growth rate

Quintiles

- > Established in 1996, Quintiles is the first global CRO in the region
- > Approximately 1,500 employees
- > Offices in 18 countries:
 - Austria, Bulgaria, Croatia, Czech Republic, Egypt, Estonia, Greece, Hungary, Israel, Latvia, Lithuania, Poland, Romania, Russia, Serbia, Slovakia, Turkey and Ukraine
- > Senior management based within each country

Assurance of Quality

- > Dedicated quality management ICH-GCP compliant
- > Clinical operations and monitoring that follow a rigorous process, optimizing delivery timelines, quality and cost-effectiveness

The Resources to Deliver on Your Objectives

Project Management

- > Highly qualified Project Managers and Clinical Trial Leaders with medical, pharmacy or life science backgrounds
- > Experience in managing global studies

Clinical Research Associates

- > Multi-skilled, multilingual, proactive CRAs and CTAs with medical, pharmacy or life sciences backgrounds
- > Wide therapeutic area experience
- > Multilingual clinical monitoring capabilities

Key Services

- > Clinical Trial Monitoring
- > Feasibility Services
- > Study Start-up Services
- > Project Management
- > Regulatory Affairs
- > Medical and Scientific Services
- > Site Coordinator Services
- > Professional Training
- > Site Contracts and Investigator Payments

Contact Us:

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