

Global Clinical Research Operations and Quality Partner

Supporting Infectious Disease, Vaccine, and Global Health Research
Across the United States, Sub-Saharan Africa, Asia, Europe, MENA and Other Emerging Markets



WHO WE ARE

Guosa Life Sciences (GLS) is a quality-driven global Contract Research Organization (CRO) delivering clinical development and operational excellence across the United States, Sub-Saharan Africa, Asia, Europe, MENA and other emerging markets.

We combine global regulatory and quality expertise with established in-country teams, site and investigator networks, and deep understanding of local languages, cultures and healthcare systems—enabling efficient, ethical, and compliant research that advances better life through better medicine.



GLOBAL QUALITY STANDARDS

International standards and best practices across the entire research lifecycle.



LOCAL INSIGHT. GLOBAL IMPACT.

In-country teams that speak the languages and understand the cultural attitudes and operational realities.



COMPLIANCE YOU CAN TRUST

Robust quality systems with a focus on ethics, safety, and data integrity.



TRUSTED PARTNER FOR GLOBAL CLINICAL RESEARCH

Supporting sponsors, CROs, academic institutions, government agencies and global health organizations through flexible and scalable partnerships.

United States
Sub-Saharan Africa

Asia
Europe & MENA

OUR GLOBAL PRESENCE



UNITED STATES

- Corporate leadership
- Quality oversight
- Sponsor engagement



SUB-SAHARAN AFRICA

- In-country teams
- Investigator networks
- Site operations
- Regulatory support



ASIA

- In-country teams
- Investigator networks
- Site operations
- Regulatory support



EUROPE & MENA

- Strategic partnerships
- Regional experts
- Study support capabilities

CORE CAPABILITIES



CLINICAL OPERATIONS

- ✓ Site identification and feasibility
- ✓ Site activation and management
- ✓ Investigator engagement
- ✓ Patient recruitment support
- ✓ Monitoring and oversight



QUALITY ASSURANCE

- ✓ Quality management systems
- ✓ Risk-based quality management
- ✓ Audit and inspection readiness
- ✓ Quality oversight throughout the study lifecycle



REGULATORY AFFAIRS

- ✓ Regulatory strategy and submissions
- ✓ Ethics committee support
- ✓ Country-specific regulatory requirements
- ✓ Compliance and reporting



PHARMACOVIGILANCE AND SAFETY

- ✓ Safety monitoring and oversight
- ✓ Adverse event management
- ✓ Signal detection and reporting
- ✓ Risk management plans



LABORATORY SERVICES

- ✓ Laboratory coordination
- ✓ Specimen management
- ✓ Sample logistics
- ✓ Central and local laboratory support



LOGISTICS AND SUPPLY CHAIN

- ✓ Study logistics management
- ✓ Cold chain management
- ✓ Importation and customs clearance
- ✓ Supply chain optimization



CONSULTING AND STRATEGIC ADVISORY

- ✓ Protocol and operational strategic planning
- ✓ Feasibility and site selection
- ✓ Market entry and setup
- ✓ Capacity building and training programs



REGULATORY AFFAIRS CONSULTING

- ✓ Regulatory pathway strategy
- ✓ Dossier preparation and review
- ✓ Authority interactions
- ✓ Product registration support
- ✓ Regulatory intelligence



IDEAL PARTNER FOR

- ✓ NIH-funded clinical research
- ✓ Infectious disease and vaccine studies
- ✓ Global health and public health programs
- ✓ Multi-country and multi-site clinical trials
- ✓ Africa, Asia, Europe, MENA and emerging market research
- ✓ Site startup and operational execution
- ✓ Quality oversight and inspection readiness
- ✓ Capacity building and training initiatives



ESTABLISHED GLOBAL PRESENCE

United States, Sub-Saharan Africa, Asia, Europe, MENA and other emerging markets



IN-COUNTRY TEAMS

Local professionals with regional expertise and deep community insight



SPEAKS THE LANGUAGES UNDERSTANDS THE CULTURES

Building trust, improving engagement, and driving successful outcomes



GLOBAL STANDARDS LOCAL EXCELLENCE

Committed to quality, compliance and integrity at every step



FLEXIBLE PARTNERSHIP SCALABLE SOLUTIONS

Your goals. Our expertise. Better outcomes together.

CONNECT WITH US

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