



A Global Full-Service CRO

Advancing human health through scientific rigor and integrity since 1977

Known for excellence in clinical trial support, we are dedicated to helping commercial, government, non-profit and academic partners achieve their biopharmaceutical development and public health goals.

With strong roots in statistical science and unwavering dedication to delivering superior research support, we have grown long, fruitful partnerships over more than 45 years.

Each of our clients experience the same high level of attentiveness and commitment, delivered by teams dedicated to making the vital contributions to science and public health that drive rapid development of patient treatments.

Partnering with
Biopharmas
Foundations
Non-profits
Government

1,500+

Professionals in
the US, Europe,
Canada and India

2,000+

Clinical Trials in
over 90 Countries

30,000+

Clinical Trial
Sites

1 Million+

Subjects Enrolled

100+

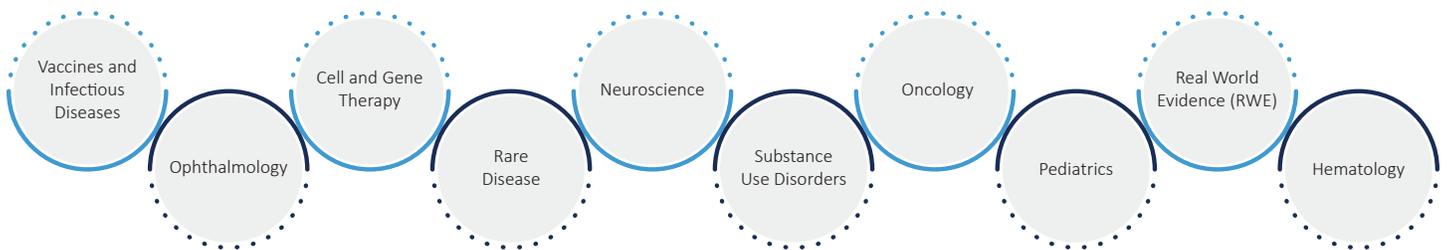
INDs Prepared
and Submitted

2,600+

Publications



Our commitment to scientific excellence, collaboration, and problem-solving helps our clients to better understand complex research problems, generate higher quality data, and make better treatment options available to patients in:



Collaborators and Consultants:

From day one, we work with you honestly and transparently and in an agile way to:

- Design studies that meet your study objectives and endpoints
- Focus on making clinical trials easier for sites and patients
- Speed your timelines with eClinical technologies, remote, risk-based, on-site study monitoring, data management and analysis excellence
- Focus on your regulatory strategies and downstream submissions
- Collaborate on publications & regulatory filings



Dedicated to integrity and scientific research

Mission

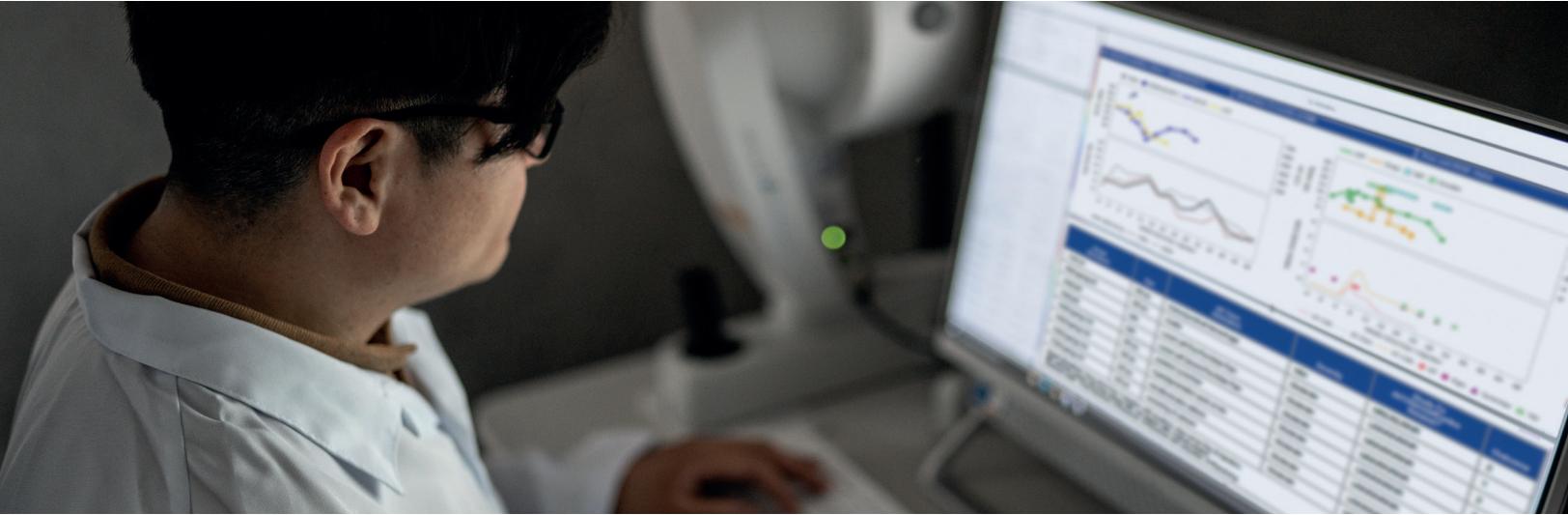
We are a global, full-service Clinical Research Organization dedicated to excellence in supporting the advancement of public health and biopharmaceutical innovation.

Vision

To be the trusted partner to clients who share our passion for improving human health in a world of ever-changing scientific research.

Values

Integrity, Agility, Passion for Excellence, Collaborative Partnerships, Intellectual Curiosity.



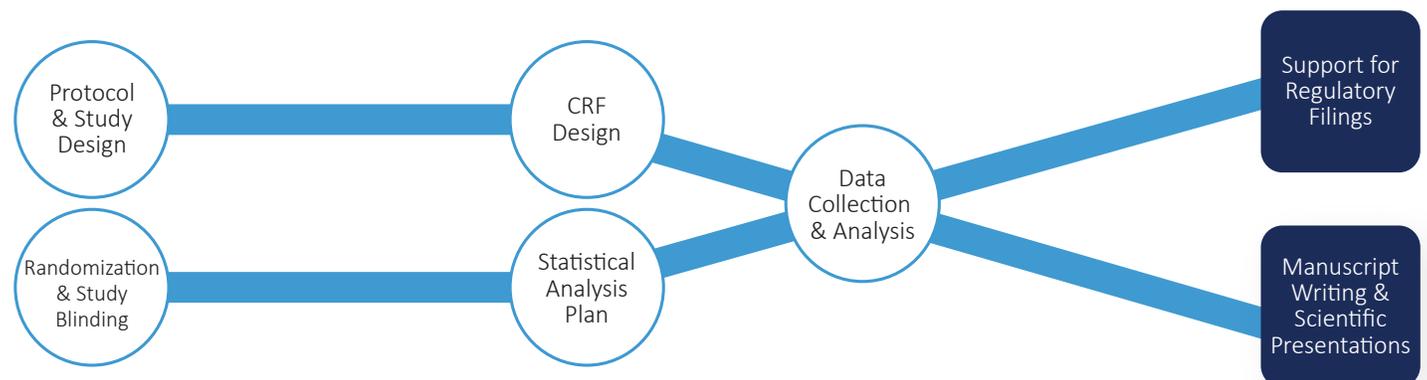
From Planning to Publication

As a full-service CRO, we can help you from the earliest stages of planning your Phase I-IV clinical trials to reporting, regulatory submissions and publications. We also have extensive experience managing registries, natural history studies and other types of clinical research, including ePRO to generate patient insights.



Biostatistical Support Across your Trial

Emmes biostatisticians get involved at the early stages of study planning, design and protocol development. To ensure data quality Emmes biostatisticians work very closely with our site monitors, medical staff, data management and IT teams throughout your clinical trial.



Experience

Our engagements with study sponsors have enabled Emmes to amass an enduring track record. We are pleased to share examples of our long-term clinical research with you.

- Age-Related Eye Disease Study 2 (AREDS2)
- AIDS Malignancy Consortium (AMC)
- Best Pharmaceuticals for Children Act (BPCA)
- Blood and Marrow Transplant Clinical Trials Network (BMT CTN)
- Clinical Research in Infectious Diseases (CRID)
- Collaborative Islet Transplant Registry (CITR)
- Consortium for Food Allergy Research (CoFAR)
- National Eye Institute Support (NEIS)
- National Institute on Drug Abuse Clinical Trial Network Clinical Coordinating Center (NIDA CTN CCC)
- National Institute on Drug Abuse Clinical Trial Network Data and Statistics Center (NIDA DSC)
- National NeuroAIDS Tissue Consortium (NNTC)
- Production Assistance for Cellular Therapies (PACT)
- Vaccine Research Center DCC (VRC)

Specialty Services

Emmes tailors its services to the demand and mission of your clinical research project. This is achieved with the following specialty services:

Data and Safety Monitoring Board (DSMB) Services

Assembled for each project, DSMBs bring an objective view on patient safety, ensuring independence from the sponsor on important decisions on trial safety.

OptymEdge

Visual Function Certification Services

Proven solutions for training and certification of Visual Acuity Examiners and Visual Acuity Rooms. We bring experience and expertise which comes from having the largest and longest standing certification organization in the industry.

» Advantage *eClinical*

Advantage eClinical Technology Suite

Emmes' unified clinical data platform that was developed by statisticians, medical officers, programmers, data managers and analysts. Our integrated ecosystem of tools, including EDC, randomization, ePRO, and sample management, allows you to design your clinical trials, ensure timely data collection, manage data quality, and report on critical outcomes.

For additional information on our services, please visit www.emmes.com