



Precision research, delivered anywhere.

Fast, agile, and affordable CRO services for clinical, nutritional, real-world evidence, and consumer research studies — managed end-to-end from protocol to final report.

78+

Studies
delivered

50+

Years'
experience

35+

Countries
covered



What we do

Clinical Trials

Full-service management of non-CTIMP trials, from protocol and ethics through to final reporting.

Real-World Evidence

Open-label trials delivered remotely using validated tools, wearables, and at-home testing.

Nutritional Research

Dietary intervention studies covering supplements, probiotics, and microbiome analysis.

Consumers / IHUT

Consumer research that validates product concepts and generates claims-ready evidence.

AREAS WE WORK IN

[GI & microbiome](#)
[Men's & women's health](#)
[Healthy ageing](#)
[Nutrition](#)
[Sports & fitness](#)

Why Comet Clinical

- ✓ Remote-first methodology reduces time and cost without cutting corners
- ✓ Experienced team — 50+ years combined delivering non-CTIMP and real-world studies
- ✓ End-to-end management from protocol design through to final report



Our electronic data capture, ePRO, and eConsent platform is provided by Trialflare, our technology partner for every remote study we run.



Ethical review for our non-CTIMP and remote studies is provided by Ethiclear — the only UK ethics committee dedicated to this kind of research.

Quality at Comet Clinical

Our own quality management system underpins every study we run, built around clear process and continuous improvement.

[Documented SOPs](#)
[Independent QA review](#)
[Root-cause CAPA process](#)
[Ongoing staff training](#)



Full lifecycle coverage, every engagement.

One partner from protocol to publication — no handoffs, no gaps, no surprises.

Protocol Design	Study design, endpoints, and statistical plan defined up front.
Ethics & Regulatory	Submissions, approvals, and ongoing compliance managed for you.
Vendor Management	Labs, couriers, wearables, translators, and other third parties coordinated.
Study Operations	Recruitment, remote monitoring, and participant support, day to day.
Data Management	Collection, cleaning, quality checks, and database lock.
Biostatistics & Analysis	Statistical analysis plan through to final results.
Reporting	Clinical study reports and sponsor-ready deliverables.
Publication Support	Manuscript drafts, congress abstracts, and dissemination.

EXAMPLE STUDIES WE SUPPORT

[Gut microbiome](#)
[Adolescent wellbeing](#)
[Male & female health](#)
[Healthy ageing](#)
[Sports & fitness nutrition](#)

[Sleep & recovery](#)

How evidence-ready is your product?

Scan for your free Evidence Readiness Score



TYPICAL INVESTMENT

£30,000 – £150,000

≈ €35,000–€175,000 · \$38,000–\$190,000

Final cost depends on study design, duration, sample size, and data collection complexity. Every engagement starts with a free scoping consultation.

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[GCP / GxP aligned](#)
[GDPR compliant](#)
[UK-based data residency](#)
[Affordable trials](#)
[Trusted experts](#)