PHARPOINT RESEARCH

SERVICE OVERVIEW

Medical Writing

High-quality, timely medical writing solutions delivered with your target audience in mind.

Our medical writing services include:

- Clinical development plans
- Investigator brochures
- Clinical study protocols
- Patients materials
- Clinical study reports
- Clinical/non-clinical studies
- Safety update reports
- Briefing documents
- FDA meeting materials
- Patient narratives

An experienced team you can trust

Our Chief Scientific Officer oversees our medical writing team, and our head medical writer has over a decade of medical writing experience across multiple therapeutic areas.

Competitive timelines & on-time delivery

Our expertise allows us to deliver high quality documents as quickly as possible. The timelines we promise, we stick to — in true PharPoint fashion.

High-quality deliverables

Quality is our number one priority. Our medical writing department is backed by a robust quality control procedure to ensure every deliverable is exactly how it should be.

PHARPOINT RESEARCH

STANDARD TIMELINES

Clinical Study Reports Delivered within 40 Business Days*

Final TLFs Received

10 BUSINESS DAYS

Initial Draft CSR Delivered

Sponsor provides consolidated comments

10 BUSINESS DAYS

Second Draft CSR Delivered

Sponsor provides consolidated comments

10 BUSINESS DAYS

CSR Ready for Sign Off/Publication

***NOTE:** Timeline assumes drafts are reviewed by Sponsor within five business days. May include roundtable meetings, if needed.*

Learn more about how PharPoint can support your upcoming study.

PharPoint Research is an award-winning and client-focused contract research organization (CRO) that offers clinical operations, data management, biostatistics, statistical programming, and strategic clinical trial consulting services to clients of all sizes.

Founded in 2007, PharPoint's team has participated in over 1,000 clinical trials with a high business repeat rate.

Speak with our team: pharpoint.com/contact