



Medical Writing Solutions From Synchrogenix

Superior writing is key to successful drug development

High quality medical writing is an essential part of any successful drug development program. Pharmaceutical and biotech companies need a trusted, agile, and experienced partner not only to write these documents, but also to act as a strategic partner and address key issues.

Synchrogenix, powered by ClinGenuity, has nurtured a powerful combination of seasoned medical writers and dedicated editors. Our rigorous, proven quality control processes ensure the highest quality of documents. Resourcing is scalable and knowledge is shared, so quality and integrity are maintained regardless of fluctuations in workload or organizational changes.

Global teams of experienced medical writers

Working with a great team is a prerequisite to success in any venture. Our team has worked in a number of therapeutic areas, including bone/muscle/joint, cardiology, dermatology, metabolic diseases, endocrinology, gastroenterology, hematology, infectious disease, medical devices, men's health, nephrology, neurology, oncology, pain/analgesia, rare disease, respiratory, vaccines, and women's health.

Our team also has experience in a wide variety of project types, including posters, presentations, abstracts, slide sets, and manuscripts describing clinical trial data, pharmacokinetics/pharmacodynamics (PK/PD) (ie, modeling, bioavailability, gender/race differences, tissue penetration), meta-analysis, data-mining, health outcomes, and health economics.

If desired, graphic support with SYSTAT SigmaPlot, GraphPad Prism, and/or Adobe® InDesign is available.

We are a highly educated workforce: over 50% of our publication writers hold doctorates, and over 80% hold advanced degrees.

Solutions from R&D through life cycle management

Our team is highly skilled in managing the transition from pre-clinical to clinical trials and development through submission and approval, to post-marketing. Our writers are fully conversant in International Committee of Medical Journal Editors guidelines. They are also adaptable to client-specific templates, guidance, and standard operating procedures.

Benefits of Partnering with Synchrogenix

- Partner with the largest consultancy of medical writers in the industry
- Leverage our proven expertise in developing global strategies to support timely and accurate release of scientific and medical information
- Allow your team to focus on data interpretation, new drug development, business development, and commercial activities—not abstract word counts, manuscript galley checks, or poster dimensions
- Increase momentum to meet required publication deadlines

We support all stages of medical and scientific information disclosure. Our writers have years of experience leading kick-off meetings, assembling drafts, coordinating author reviews, managing journal and conference submissions, and performing galley checks.

Powerfully communicate the science of drug development

With almost 30 years of writing and document support experience, Synchrogenix has honed a unique methodology and built a team of writers who are leaders in their field. With flexible and scalable solutions, expertise across the entire drug development life cycle, Synchrogenix offers our clients a complete outsourcing capability for all of their medical writing needs.

About Certara

Certara is a leading provider of decision support technology and consulting services for optimizing drug development and improving health outcomes. Certara's solutions, which span the drug development and patient care lifecycle, help increase the probability of regulatory and commercial success by using the most scientifically advanced modeling and simulation technologies and regulatory strategies. Its clients include hundreds of global biopharmaceutical companies, leading academic institutions and key regulatory agencies.

For more information visit www.certara.com or email sales@certara.com.