DiagnoSearch

Scientific Writing

Clear, Compliant, Submission-Ready Documentation

Our dedicated team of physicians, pharmacists, and scientists deliver end-to-end Regulatory and Scientific Writing Services that span the entire clinical development lifecycle from strategy to submission. We specialize in producing high-impact, transparent, detail-oriented documents to accelerate regulatory approvals and journal submissions.

100+
Submissions

30+
Years Experience

10+
Therapeutic Areas

Services Offered

- Concept Documents
- Clinical Development Plan
- Clinical Trial Documents (Phases I–IV) to name protocol synopsis, protocol and amendments, consent / assent forms, IB, CSR and others
- Educational Materials
- Regulatory Submissions / Electronic Submission Documents / eCTD Documents
- Pharmacovigilance / Safety Narratives / Reports & Risk Management Documentation
- Real-World Evidence (RWE) & Health Economics (HEOR)
- Scientific Publications & Medical Affairs Assets
- Labelling Documents (Pack Inserts, Product Monographs etc.) and CCDS

The DiagnoSearch Advantage

- Proven track record of contributing to 100+ successful regulatory submissions with minimal regulatory queries or revisions
- Publications in peer-reviewed journals and systematic reviews
- Documentation tailored for strategic impact, regulatory speed, and clarity
- Expertise across 10+ therapeutic areas including Vaccines, Oncology, Rheumatology, Immunology, Infectious Diseases, Metabolic Disorders, and Rare Diseases
- Trusted Medical and Scientific Writing Solutions tailored for various geographies

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