

Senior Medical Writer 1

Job ID
REQ-10037230
Ene 26, 2025
India

Resumen

To write, review and manage the preparation of high quality clinical documents for CPO's and global organization. Provide authoritative documentation related consultancy to other line functions.

About the Role

Senior Medical Writer I

Location – Hyderabad #LI Hybrid

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To write, review and manage the preparation of high quality clinical documents for CPO's and global organization. Provide authoritative documentation related consultancy to other line functions.

Key Responsibilities:

- To author, review and independently manage high quality clinical documents: Clinical Study Reports (CSR) including narratives, Protocol, Informed Consent Form (ICF).
- To write CTD modules and other safety documents (DSURs, RMPs) independently
- Liaise with medical/clinical experts, statisticians, investigators in concept development when protocol is being developed and work in a collaborative fashion for global/CPOs
- Contribute to planning of data analyses and presentation to be used in CSRs
- Ensure compliance of documentation to internal company standards and external regulatory guidelines.
- Supervise outsourcing to external medical writers, if required.
- Training and mentoring of associates as required.
- Contribute to cross-functional communication to optimize feedback and input towards high quality documents. Maintain audit, SOP and training compliance.

Commitment to Diversity & Inclusion: :

We are committed to building an outstanding, inclusive work environment and diverse teams representative of the patients and communities we serve.

Essential Requirements:

- Minimum university life science degree or equivalent is required. Advanced degree or equivalent education/degree in life sciences/medicine/pharmacy is desirable.
- 3 years of regulatory medical writing experience or other relevant pharma industry experience combined

with scientific and regulatory knowledge, plus in-depth knowledge of medical writing processes.

- Good communication skills (written, verbal, presentations)
- Good operational knowledge of clinical trial reporting.
- Good knowledge of biostatistics principles.
- Strong ability to prioritize and manage multiple demands and projects.
- Knowledge of and experience in global regulatory environment and processes (key regulatory bodies, key documents, approval processes, safety reporting requirements).
- Good experience in managing global, cross-functional teams or simple global projects.

Desirable Requirements:

- Demonstrated ability to establish effective working relationship in a matrix and multicultural environment.
- Demonstrated presentation and diplomacy skills.
- Strong customer-oriented mindset.

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Benefits and Rewards: Read our handbook to learn about all the ways we'll help you thrive personally and professionally: <https://www.novartis.com/careers/benefits-rewards>

División

Operations

Business Unit

Innovative Medicines

Ubicación

India

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Hyderabad (Office)
Company / Legal Entity
IN10 (FCRS = IN010) Novartis Healthcare Private Limited
Functional Area
Research & Development
Job Type
Full time
Employment Type
Regular
Shift Work
No
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Accessibility and accommodation

Novartis is committed to working with and providing reasonable accommodation to individuals with disabilities. If, because of a medical condition or disability, you need a reasonable accommodation for any part of the recruitment process, or in order to perform the essential functions of a position, please send an e-mail to diversityandincl.india@novartis.com and let us know the nature of your request and your contact information. Please include the job requisition number in your message.

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