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Senior Medical Writer

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Location: Belgium

Category: arts-design-entertainment-sports-and-media

As a Senior Medical Writer you will be joining the world’s largest & most comprehensive clinical research organisation, powered by healthcare intelligence.
As a Senior Medical Writer you will be supporting a Top-5 pharma company. You have the chance to help our customer to deliver best-in-class regulatory and submission documents. While being stably employed with ICON, you will be fully embedded with our client.

Responsibilities:
Independently produces clinical and regulatory documents in collaboration with Principal Medical Writers . These documents may include clinical study level documents such as Clinical Study Protocols or Clinical Study Reports. This includes Phase 1 in patients to Phase 3 across multiple therapeutic areas.

Thoroughly analyze and critically interpret data to determine the best approach to composing each document, applying lean writing strategies.

Acts as the primary contact for the study team in relation to the preparation and timelines (incl. planning) of assigned documents.

Facilitates the review of the documents and ensures that documents are submission-ready and finally approved. When preparing documents you will be directly in contact with QC personnel and publishing specialists who support you in document preparation.

Minimum requirement of a bachelor’s degree; Master's degree preferred

3-5 years Medical Writing experience or other relevant pharmaceutical experience combined with scientific and regulatory knowledge

Strong writing skills and the ability to convert scientific data into a clear, scientifically sound, well-structured messages.

Proficient in independently writing several types of clinical/regulatory documents (mainly CSRs, Protocols) including leading creation, coordination of the authoring functions, facilitation of the review of the documents and ensuring that documents are submission-ready and finally approved.

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