

Associate Medical Writer

Role

As a Medical Writer with Azur Health Science you will assist our clients with the development of high-quality medical document deliverables for the preparation of regulatory and other medical communication content in accordance with applicable internal, external, and international guidelines.

Teamship

As a Medical Writer you are the focal point of a cross-functional project team that has the combined aim to produce a high quality document to inform a specific target audience. This target audience could be clinical study teams, ethics committees, regulatory agencies, clinical trial patients, or scientific professionals in the field. This is an important role in the process of bringing novel products to market and monitoring the safety and effectiveness of marketed products in the real world. The medical writer's role requires proactive diplomacy to manage the requirements of the project team as well as to manage the project itself.

Responsibilities and Accountabilities

Working with Azur, you will be required to actively contribute to the development of our team based on the foundational ethics and values of the company. Azur was founded on the principle of Teamship where there is a shared accountability and where everyone has an equal voice.

As a member of the Azur team, we expect you to respect all individuals of the team and to celebrate successes together with us, as well as being willing to share from your mistakes, in order that we might learn as a team.

The Azur team provides your learning and support network. While we expect you to take ownership and accountability of your projects, you must never work in isolation, and you must communicate regularly with the Azur team about the progress of your project. Your ultimate accountability lies with Azur Health Science.

Required Skills

As you work for Azur, you will need to develop the following skills. Those that were assessed during the recruitment process, we expect you to work on, develop, and deepen during as your career with us develops and matures.

- Ability to produce clear, concise scientific writing in English
- Ability to follow and work according to the guidance and regulations of the industry, such as Good Clinical Practice and data confidentiality and privacy regulations
- Ability to synthesise data, and non-clinical and clinical concepts from a broad range of disease areas
- Ability to form strong collaborative relationships while working effectively in cross-functional, remote, geographically and culturally diverse teams
- Strong project management skills to run and organise project teams empowering them to produce multiple tasks within agreed timelines
- Strong interpersonal, verbal, and written communication skills
- Ability to think through and work around problems logically in response to changing conditions

Career Progression

The usual career progression is as follows:

Role Title	Expected Timeframe in Role
Associate Medical Writer	1-2 years
Medical Writer	2-4 years
Senior Medical Writer	2-4 years
Principal Medical Writer	2+

If you are interested in this position, please [complete the form](#) on the recruitment page.