This position "Medical Writer - Senior" covers medical writing activities across all phases of development for small molecules, biologicals, cell based and modern targeted/personalized therapies. Experience in any of these areas is a prerequisite.

Tasks & Responsibilities:

- Authoring high-quality regulatory documents
- Such as IB, CTD summary documents (2.6, 2.7) and overviews (2.4, 2.5)
- Orphan Drug applications, RMPs, and PIPs
- Compilation of any documentation required for approval and conduct of clinical studies. Documentation to support agency meetings (briefing books, presentations)

Requirements:

- Advanced degree in life sciences, medicine or veterinary medicine
- Broad experience in medical writing (>5 years) on wide range of regulatory documents
- Understanding of FDA and ICH regulations and guidelines
- Ability to work independent, analyse and summarise data (clinical, non-clinical, quality)
- Ability to manage multiple tasks simultaneously under time constraints
- Excellent communication skills in English and a second language, preferably German
- Strong knowledge of Microsoft Office and graphical software

We offer:

A professional and friendly working environment located in Munich. Working together with a dedicated small and effective team, we provide an outstanding and more than competitive salary, with a very attractive bonus system tailored to your expertise and our business success.

Contact:

If you are interested in working in an innovative and team-oriented environment, please send your application documents to Dr. Stefan Blesse, blesse@granzer.biz .

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