Biomapas is a functional and full outsourcing solution provider to the global life science industry, with key expertise in Clinical Trials, Regulatory Affairs and Pharmacovigilance. With headquarters in Lithuania and offices in Switzerland, Russia, Georgia, Ukraine, Poland, Kazakhstan and Sweden, Biomapas operations are spread over 5 continents, concentrated in Europe, Russia and former CIS region.

Biomapas is looking for a **Quality Assurance Director** to join leading position and strengthen our further development and expansion of overall company activities globally.

Role responsibilities:

- Develop and implement Company quality assurance and management strategy according to the best global standards;
- Oversee all quality assurance efforts within organization;
- Maintain and improve quality management system;
- Ensure GCP audit revenues plan execution;
- Consult Company employees regarding all quality related questions;
- Ensure that Clinical Operations, Pharmacovigilance, Regulatory Affairs processes conform to the applicable regulations, GCP, GVP and other applicable practice guidelines, ISO 9001 and ISO13485 standards requirements;
- Define and ensure implementation of Quality Assurance Department objectives;
- Establish and implement quality and performance metrics and analysis methods in collaboration with heads of departments;
- Oversee and ensure compliance with QMS documentation and local and international requirements and standards;
- Host second, third party audits or regulatory inspections in the company and oversee the implementation of corrective and preventive actions plans after the audits;
- Participate in the training process of the staff by preparing and presenting quality and legal requirements relevant trainings.

Role requirements:

- 10 years of relevant executive quality management experience in B2B environment;
- Minimum 5 years of experience in managing and leading teams;
- Experience in strategic development and execution of quality management system;
- Experience in a Clinical, Regulatory Affairs or Pharmacovigilance service provider environment;
- Knowledge of GCP, GVP, ISO 9001 and ISO 13485 standards;
- Experience in operations with any electronic QMS;
- Ability to lead/supervise activities in a high-pressure, fast paced and changing environment to ensure objectives are met in a timely manner;
- Fluent English language.

In Biomapas you will find supportive work environment with guarantee for professional and personal development, as well as competitive salary and benefits and many more initiatives that will make your daily office life comfortable.

Please apply to cv@biomapas.com

Be kindly informed that only selected candidates will be contacted.