BIOMAPAS
Clinical, Regulatory & Pharmacovigilance

We keep promises. Always.

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 an **International QPPV** to oversee a global network of local and regional QPPVs and represent the QPPV Office in the Pharmacovigilance leadership team.

Key responsibilities include:

- Line management of a team of EU QPPVs, QP Office Scientists and QP Office associates;
- Represent the QPPV Office in the PV leadership team;
- Oversee the global network of QPPVs, including dotted line management of Regional QPPVs;
- Act as the appointed EU QPPV or deputy EU QPPV for the Customer in accordance with a separate QPPV Job Description;
- Management of projects with the scope of the EU-QPPV role and / or the management of a full pharmacovigilance system / parts of the global pharmacovigilance system;
- Ensure that all pharmacovigilance activities are in compliance with Biomapas and Biomapas contractual partners' procedures;
- Continuously develop his/her professional and personal skills and participating in pharmacovigilance relevant trainings delivered by Biomapas and/or Biomapas contractual partners;
- Supporting Biomapas business development department in promotional activities related to QPPV function, generating answers to pharmacovigilance RFIs and RFP documents;
- Have oversight on Safety Database all other IT systems/tool used by the PV department, managed within the QPPV Office.

Key requirements for the role:

- University degree in Life Sciences, preferably master or higher degree;
- In–depth, demonstrated experience in most areas pertaining to pharmacovigilance (including QPPV responsibilities) and pharmacovigilance legislation; at least 5 years of work experience in PV functions;
- At least 3 years line management experience;
- Excellent knowledge of international regulations (EU GVP Modules, ICH, FDA);
- Strong computer literacy with intensive experience with safety databases;
- Ability to set up and implement PV strategies, develop PV systems and procedures, and elevate and resolve issues;
- Ability to accurately and effectively evaluate, interpret and present medicine safety data and processes;
- Professional level in English (both oral and written);
- Strong communication and relationship building skills;
- Outstanding analytical skills;
- Proactivity, organizational excellence and project management skills;
- Strong organization and time management skills;
- Ability to plan and prioritize workload (own and team);
- Ability to interpret and apply global medicine safety regulations.

Please apply to hr@biomapas.com

In case you would like to get more information – please do not hesitate to contact **Raimonda Klimienė**, HR and Training manager by **+370 69 815 736**.

Be kindly informed that only selected candidates will be contacted.