# **Belgium Jobs Expertini**®

## Real World Evidence Consultant (candidate pool)

# **Apply Now**

Company: Keyrus Life Science Location: Belgium Category: computer-and-mathematical

## **Job Description**

Keyrus Life Science is looking on a regular basis for **Real World Evidence Consultants** to join our consulting team for clients (from big pharma to small biotech) based in **Belgium** 

This job description is an example on how your job could look like. Once we have a specific project and client you will be informed and you will have of course the choice to be presented at our client.

Your potential challenges could be:

Support Statistical programming activities in Real World Evidence.

Provide input on study design and analysis plans where necessary.

Review the study concept, protocol design, analysis plans and other study-related documents.

Evaluate the quality of the database.

Develop, validate, maintain, and document analysis programs, tables, charts and associated metadata following standard working practices.

Ensure that statistical outputs are produced in an efficient manner.

#### Profile

5+ years experience in SAS programming.

2+ years experience in Real World Evidence, Epidemiology, or Health Outcomes environment.

2+ years experience with large health care databases such as Clinical Practice Research Database (CPRD) or IBM MarketScan.

Advanced statistical programming skills in SAS o solve complex programming tasks.

Good **study design and epidemiological/statistical analysis experience**, including understanding of drug development, observational and epidemiological endpoints and assessments.

Knowledge of matching algorithms such as propensity score matching is a plus.

Knowledge of R is a plus.

Good written and verbal communication skills. Experience in communicating with all levels of personnel across various departments, in domestic and international environments.

Good command of English, both verbal and written.

Demonstrated experience in participating in collaborative work teams at local and global levels.

Good project management skills including experience in managing multiple tasks concurrently.

Ability to work under pressure and without close supervision.

#### What we offer

Joining our consulting team means giving your career a step forward. Thanks to the privileged partnership we have with our well-established clients, leaders in their domain, we offer our consultants unique projects to develop their skills, and put their professional experience in the pharma industry on the fast track.

As an expert on the client side, you can count on us to support you at each step of your project thanks to regular touch points with our Key Account Manager. We then also carefully choose with you what would be your next assignment that is matching your career goal whether it is in Consulting or in one of our Functional Services Platforms.

We invest considerable time and resources in training our staffitechnical and non-technical

courses) to continuously support you in your personal and professional growth. On top of this, you will receive a **complete salary package including attractive extra-legal benefits** (company car, petrol card, meal vouchers, group insurance, hospitalization...) based on your background and experience.

#### Who we are

Keyrus Life Science is an international Consulting, Contract Research Organisation and Functional Services provider with a reputation for transparency and integrity, highly focused on being able to deliver with excellence. Our 'human' approach to service provision is what differentiates us from our competitors. Combined with our high caliber staff, this approach has allowed us to become a key player in clinical research.

At Keyrus Life Science we're proud of our commitment to delivering services of the highest quality, not only skilfully, efficiently and reliably, but also with sincerity and genuine care for our clients' projects, priorities and reputation. Thus we maintain and advance our vision and our standards, all the while ensuring that we serve your interests better.

From early- to late-stage drug development, our range of services includes: Project Management, Clinical Operations, Pharmacovigilance, Quality Assurance (GxP), Regulatory Affairs Strategy & Support, Data Management, Medical Review & Coding, Biostatistics, Medical Writing & Medical Information, Real World evidence services.

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