[노보노디스크제약] Real World Evidence Associate (1 년 계약직)

<Job Description>

Contract Period
The contract period is 1 year, and this position can be possibly extended in its contract after 1 year.

Position's key accountabilities

- Support execution and implementation of non-interventional studies and other Real-World Evidence generating studies to support the NNPK evidence generation plan with partners and with vendors.
- Contribute to driving innovation in RWE analytics by implementing new analytic methods, study design and technologies to answer research and business questions.
- Support conducting RWE studies using internal and external real world data to generate unique disease insights and epidemiological information to improve patient care and support business needs.
- Communicate RWE matters with key internal stakeholders and scientific community.

Nature and scope of Main accountabilities

- Collaborate with external vendors on RWE projects and manage communication, contracting and project follow up with external vendors (ex. CROs) and ensure expected deliverables and meet timelines / challenges on content for low or medium complexity projects.
- Identify Real World Data sources (electronic medical records, registry database, patient reported outcomes survey, and national claims database) which best support business needs through literature reviews and desk research etc.
- Able to interpret, understand and discuss data results, protocol, SAP, NSR and other study related materials, and collaborate with internal stakeholders and external vendors with timely and efficient communications.
- Collaboration in study operations with RWE manager:

  - Provide input for feasibility and operation perspectives to study outline,
  - Prepare study protocol, informed consent and amendments, if applicable,
  - Provide input to study and project related documents (e.g. CRF, NSR, Study Validation Plan),
  - Ensure availability of other study related documents, e.g monitoring guidelines,
  - Plan and conduct investigator meetings and other meetings incl. Investigator Result meeting(s),
  - Ensure audit and inspection readiness,
  - Handling protocol deviations (if applicable) to ensure high compliance,
  - Ensure compliance with all existing applicable requirements, and overall ensuring compliance with GCP and other applicable external regulations for the non-interventional studies as well as Criteria for medical drug promotion and local regulations in the local meetings within and outside the study community.

- Background knowledge of industry trends and best practices specifically related to epidemiology/outcomes research.
- Good knowledge and understanding of applicable regulations in pharmacoepidemiology, pharmacovigilance and clinical development.
- Conduct literature reviews and online research to support Real World Evidence projects.
- Combination of strong methodological quantitative knowledge, strategic, innovative thinking, and communication skills.
- Prepare presentation materials (in PowerPoint) and assist in document development (in Word & Excel) for Real World Evidence protocols, reports, manuscripts, posters, etc.

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- Degree in statistics, public health, or other health related fields.
- Master's degree in a field such as epidemiology, biostatistics, statistics, bioinformatics, economics or similar is preferred.

- Minimum of two to three years of related experience conducting research in the pharmaceutical industry, contract research organization, healthcare provider / HTA or academic institute; or experience in a closely related discipline within the pharmaceutical industry.
- The ability to develop understanding of retrospective and prospective research methods, late phase study designs, HEOR and related methodology.
- Record of publications in observational research is a plus.
- Experience working with longitudinal RWD (e.g., claims, electronic health records, and/or surveys).
- Experience in the conduct of non-interventional study, late phase study, or database study.
- Experience in using statistical software (e.g., R, SAS, Python, SPSS, and/or Stata).
- Collaborative approach to problem-solving and ability to plan and multi-task.
- Excellent written and oral communication skills.
- Excellent communication, presentation, interpersonal skills, both written and spoken.
- Proficiency in Excel, Word and PowerPoint.
- Fluency in written and spoken English.

<고용형태> 계약직

<전형방법> 1 차 서류전형 - 2 차 면접전형

<접수방법> 홈페이지 접수 https://careers.novonordisk.com/job/Seoul-Real-World-Evidence-Associate-Seou/794804401/

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