reMYND is a biopharmaceutical company focused on the development of treatments against protein-misfolding disorders, with two business units:

The Drug Discovery and Development (DDD) unit, driving our own pipeline of disease-modifying treatments against diabetes, Alzheimer's, Parkinson's and orphan diseases

The CRO unit, a world-wide leader of preclinical Alzheimer's disease contract research in our proprietary transgenic mouse models

Open position: Clinical Operations Associate

The Clinical Operations Associate provides management of the clinical operations activities ensuring GCP-compliant, timely and within budget conduct of reMYND-sponsored clinical trials.
The ideal candidate:

- Tracks the execution of the operational strategy for all clinical programs in line with project deliverables and according to GCP and other applicable regulatory guidelines
- Follows up that clinical deliverables are met on time, within budget, and according to quality standards
- Monitors that clinical studies in all development Phases are being conducted in line with the protocol, standard operating procedures (SOPs), good clinical practice (GCP), and applicable regulatory requirements
- Supports monitoring visits for assigned protocols and study sites (also virtually if required) in several therapeutic areas, and identifies and escalates quality issues observed to the CMO
- Contributes to development and review of (clinical development plans and) clinical trial documents
- Ensures optimal use of resources (internal as well as external)
- Supports the CMO with appropriate reports and communication about study progress, site management issues, and monitoring visit findings including suggestions for problem solving
- Liaises with external clinical operations team and resources
- Ensures development of and maintains oversight over SOPs for clinical operations
- Is accountable for clinical operations aspects during future Regulatory Agency inspections and internal audits
- Reports to CMO

Specific expertise and qualifications:

- Operational experience with multiple clinical studies in several phases
- Solid knowledge and experience with GCP and clinical trial management
- Able to function well in a matrix structure and organization which collaborates with external partners
- Focus on execution and deliverables - organized, quality driven and timely
- University Degree in (bio)medical, paramedical, pharmaceutical sciences, or a qualified degree in nursing
- Willing to travel (if the Covid19 situation allows)

Please send your resume and motivation letter to applications@remynd.com, mentioning "application Clinical Operations Associate"