LINK Medical Research is an European, full-service contract research organization (CRO) that provides services within clinical trials, regulatory affairs, pharmacovigilance, health economics and quality assurance for the pharmaceutical industry and producers of medical devices. LINK Medical was established in 1995 and we are currently more than 100 employees based in Oslo, Stockholm, Malmö, Copenhagen and Berlin. Our mission is to improve and accelerate our customers 'projects development.

Senior Regulatory Manager

LINK Medical Research is looking for highly motivated Senior Regulatory Manager to join our Regulatory group. LINK Medical is growing and the portfolio of cross-functional projects is increasing. As a Senior Regulatory Manager, you will work in close collaboration with members of several project teams and members of the international Regulatory group. You may also be outsourced to clients in a close proximity to your home. In this position you will also have the opportunity to mentor junior members of the Regulatory team.

Job Profile

Qualifications

- ☐ Basic education within relevant field, equivalent to degree from University or University College.
- ☐ Good knowledge of Regulatory Affairs.
- ☐ Ability to show initiative.
- ☐ Ability to work independently.
- ☐ Good knowledge of several therapy areas.
- ☐ Ability and desire to fulfil our customer promises: Quality, punctuality and communication.
- ☐ Fluent in Swedish and English (written and oral).
- ☐ Basic IT knowledge.

Position objectives

- ☐ Complete projects/services according to the agreement with project manager, line manager and customer.
- ☐ Reporting implementation and progress of projects and other services according to agreement with project manager, line manager and customer.
- ☐ Behavior according to our promises (quality, punctuality, communication) and values (credibility, focus on solutions, job satisfaction).
- ☐ Contribute to the development of their area of expertise within LINK Medical.

Main responsibilities

- ☐ Completing task(s) according to applicable regulations, agreements and budget (e.g. maintenance of marketing authorization, strategy/development work, and regulatory intelligence).
- ☐ Ensuring that defined objectives and plans are met and complied to.
- ☐ Ensuring customers' positive perception of the collaboration with LINK Medical.
- ☐ Active project participation
- ☐ Cooperation with colleagues in LINK Medical.
- ☐ General supervising within relevant fields.

- ☐ Follow current administrative routines.
- ☐ Other relevant tasks given by the line manager.
- ☐ Participate in relevant courses, conferences, symposia and meetings as agreed with Line Manager/Customer.
- ☐ Develop training material and conduct internal and/or external courses/training.

LINK Medical offers:

LINK Medical offers an exciting and challenging position in a European CRO that has a strong local presence. The company focus on collaboration, sharing of experience and continuous development. You will be given meaningful tasks requiring a good collaboration between industry, clinical and scientific teams and the authorities. We offer a competitive salary, pension and insurance scheme and a bonus scheme for all employees.

The position is located in Stockholm.

If you have any questions about this position please contact Managing Director in Sweden, Jan Hellqvist (+46 70 913 51 25)

Please send your application and CV to;
Jan Hellqvist, jan@linkmedical.se as soon as possible.

All applications are treated confidentially.