



It's all about trust

TFS

TFS, with headquarters in Lund, Sweden, is the largest non-listed European clinical Contract Research Organization (CRO). Clinical services are provided by four business areas; TFS Explore™, TFS Develop™, TFS People™ and TFS Academy™, with annual net revenue greater than € 40 million, global operations in 21 countries, and 500 employees worldwide. Detailed information about TFS business areas, global locations and recent press releases can be obtained at www.tfscro.com

Regulatory Affairs Manager

TFS is now looking for a Regulatory Affairs Manager who appreciates a flexible working environment filled with interesting and stimulating assignments.

TFS is working with both international and national projects within many therapeutic fields. The Regulatory Affairs department is involved in plenty of exciting projects in various phases, such as registration, certification strategy, product planning, in early development phase, clinical trial application and marketing approval as well as maintenance after approval.

Our client base is growing in the Copenhagen and Malmö/Lund area and we are now looking for a well-experienced Regulatory Affairs Managers who, like ourselves, appreciates a flexible working environment filled with interesting and stimulating assignments and who enjoys working with highly competent colleagues in a smaller but dynamic group.

The main responsibilities of the Regulatory Affairs Manager will include:

- Preparation, review and compilation of regulatory submissions
- Liaison with authorities/notified bodies
- Advice/answers to regulatory issues/questions
- Maintain a high level of competence regarding international regulatory requirements and regulatory procedures

Requirements/Qualifications:

- Suitable academic education in life science such as pharmaceutical science, medicine, biomedicine, biology or similar
- At least 5 years experience from working with regulatory affairs within the pharmaceutical industry or regulatory authorities. Experience from Medical Device and/or In vitro Diagnostics is an advantage.
- Good communication skills in spoken and written Swedish and English
- Risk Management file and Claims documents
- Biological/Clinical Evaluation





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TFS Trial Form Support will offer:

A dynamic and growth-oriented organization with a work environment distinguished by professionalism, integrity and responsibility.

Welcome to join us and to contribute in placing TFS Trial Form Support among the top 10 worldwide CROs.

Your job will make a difference!

For more information about this position please contact:

Tina Thuesen, Unit Manager, Denmark, at +45 (45) 65 20 68 or e-mail tina.thuesen@tfscro.com

Cecilia Falkenberg, Director Global Medical Affairs at +46 (46) 280 18 06 or e-mail cecilia.falkenberg@trialformsupport.com

We are looking forward to receiving your application with included CV and a personal letter. We practice a continuous selection procedure, so please submit your application via this link <http://www.webforum.com/form/tfsjobb/form.asp?sid=591893229> or click on "Submit Your Application" on our website as soon as possible, however no later than the 1st of February 2012.

