Full Time Regulatory Affairs Consultant

Orphix is a highly successful medicinal product regulatory consulting organisation located in Munich.

Due to an ever-increasing portfolio of client companies based in the USA and Europe we are seeking a highly experienced and scientifically competent regulatory affairs consultant who would like to undertake interesting and challenging work on a full-time basis (40 hours per week).

Candidates must have at least 4 years European and US regulatory experience including; CHMP scientific advice procedures, FDA Type B and C meetings, paediatric investigational plans, orphan drug applications, clinical trial applications, INDs and marketing authorisation applications.

The ideal candidate would work independently, produce written documents of a high standard both in terms of presentation and scientific content, and be able to interact directly with client companies confidently, knowledgeably, and diplomatically.

Fluency in written English is essential.

Orphix welcomes applications from talented people with outstanding expertise in fields within our core strategic interests. Interested in being part of a motivated, interdisciplinary, and international team? Please send your CV via our website www.orphix.com!

Dated 26 February 2021