PhaRA

PHARMACEUTICAL REGULATORY AFFAIRS

PhaRA is a consultancy firm in EU and BENELUX regulatory affairs, located in Antwerp (10 min. walking distance from central station)

We provide strategic and hands-on regulatory support in various therapeutic areas and ranging from life-threatening diseases to OTC and borderline products

Regulatory Affairs Manager (m/f)

JOB DESCRIPTION

Depending on your level of experience // Responsible for, or assist with, the preparation, submission, and maintenance of Marketing Authorisations and Clinical Trial Applications, updating national labeling // Agency interactions in BENELUX countries and with the EMA // Lead or provide assistance for regulatory projects at EU level, such as the preparation of scientific advice, paediatric plans, management of pre- and post-marketing activities for EU procedures // Gain exposure in global development activities

We offer a challenging job with a high level of responsibility, flexibility and independency. You will receive support from an enthusiastic team. We try to create a stimulating environment and an attractive workspace

PROFILE

Masters in (pharmaceutical) sciences or PhD // Preferably a few years of industry experience (regulatory affairs) // Accuracy is your second nature // A dynamic person who can operate in a small and flexible working environment // Good project manager and team player // Willing to attend national and international conferences and courses

If you are looking for a new challenge, contact info@phara-consulting.be