

Your Specialties, Our Expertise

Biocides Regulatory Affairs Project Manager (F/M)

| Contract | Speciality | Start date | Location |
|-----------|------------|------------|----------------------|
| Permanent | Biocides | ASAP | UK – Germany - Spain |

Are you ready to take your career in Regulatory Affairs to the next level? Your expertise will shape the future of compliance and innovation in the biocides sector.

Why join CEHTRA ?



Job main purpose

Accelerate and diversify the development of CEHTRA in the Biocides regulations, in coordination with the Biocides market leader. Take advantage of CEHTRA's innovative digital tools to offer differentiating offers to customers.

CEHTRA is a social economy company and the French leader in regulatory, toxicological and ecotoxicological consultancy. Our reason for being : The prevention of toxicological risks for humans and the preservation of biodiversity.

CEHTRA offers rich and diversified career paths in expertise and management.

CEHTRA is part of the H2B Group, a forerunner in conformity assessment for climate, health and the environment.

CEHTRA is present in the world's most dynamic product registration markets: chemicals, cosmetics, biocides, plant protection, pharmaceuticals, veterinary, packaging, medical devices, etc.

CEHTRA is headquartered near Bordeaux, with offices in France (Lyon, Paris), Germany, Spain, the UK, Canada and India.

Compensation

Receive a salary tailored to your expertise and qualifications.

Key missions

- Manage biocides products dossiers, product family dossiers, and active substance dossiers to ensure compliance with European Biocides Regulation (BPR) or global legislations
- Build and maintain strong relationships with customers, delivering tailored regulatory strategies and advice
- Contribute your knowledge of biocides regulations in regions like the US, Latin America, or Asia, adding global value to our offerings.
- Be part of a team of multidisciplinary experts while enjoying the autonomy to manage regulatory projects effectively

Your profil

| Academics | Experience | |
|--|--|--|
| Holds a degree in chemistry, microbiology, or a related scientific discipline. | Brings a minimum of 5 years of experience in a similar regulatory affairs role. | |
| Skills | Team spirit | |
| Confident in customer relationship | Team settings Be proactive Take on challenges Organizational skills Autonomous | |
| English fluent (French is a plus but not essential) | | |
| Project management | | |

Position open to people with disabilities: flexible recruitment process (videoconferencing), teleworking possible and flexible working hours.

Interested?

Please send your CV and cover letter to:



