



Your Specialties, Our Expertise

REGULATORY TOXICOLOGIST

Contract	Specialty	Start date	Location
Full-time	Toxicology	ASAP	Delhi, INDIA

**Have passion for biodiversity and environment?
Broaden your horizons and enrich your career with CEHTRA!**

Why join CEHTRA?



CEHTRA is a social economy company, the French leader and #3 worldwide in technical assistance and regulatory, toxicological and ecotoxicological advice for the chemical industry. Our ambition : to promote biodiversity by responding to the strategic challenges of our clients.

CEHTRA offers to talented scientists rich and diverse career paths to expertise and management.

CEHTRA is part of the H2B Group, specialised in testing, inspection and certification related to health and environmental issues, on a human scale and with a strategic focus on digital transition.

CEHTRA is active on the most dynamic Life Science markets: Chemicals worldwide registration, Plant Protection Products, Biocides, Cosmetics, Pharma, Packaging, Medical Device...

Headquartered near Bordeaux, CEHTRA has a European/global footprint with 50 consultants in France (Bordeaux, Lyon, Paris), Germany, Belgium, Spain, United Kingdom, Canada and India.

Job main purpose

Play a central role in human health risk assessment (with some brief involved in physicochemical and environmental risk assessment) for chemicals, biocides or cosmetics.

Key Missions:

- Write toxicological monographs for pharmaceuticals, safety assessment reports for cosmetics and prepare REACH dossiers.
- Conduct literature search to retrieve data on human health or environmental toxicity endpoints.
- Assist in the development of our services to International clients which includes calling, interacting, and visiting client locations in India.
- For this job, you must have a strong understanding of toxicological endpoints, human health risk assessments (PDE/MoS calculations) and OECD or related guidelines.
- Training for the role will be provided by people in India or in Europe depending on the mission.

Your profile:

Academics

Master's degree in Toxicology / Pharmacology with 2-5 years of experience in the related field.

Experience

Knowledge of European and/or Indian Regulations and experience of liaison with regulatory authorities (BIS, CDSCO, CIB, etc.) will be an added advantage.

Skills

- English, writes business letters and reports, and possesses good conversational / telephone skills
- Proficiency with MS Office

Team spirit

- Self-motivated and ability to work remotely with international consultants
- Strong work ethic & professionalism

Compensation

Attractive, depending on the breadth of your experience

Interested?

**Please send your CV and cover letter to:
recruitment@cehtra.com**

www.cehtra.com