Junior Pharmaceutical Validation Consultant

Técnicas de Diseño y Validación, SL

Job description

Consulting company TDV SL is looking for a Junior Pharmaceutical Validation Consultant to join Barcelona team and provide highly qualified technical and scientific services to the Pharmaceutical Industry.

Functions

Reporting directly to Project Managers his/her role and responsibilities will include:

- Collaboration in preparation of regulatory affairs documentation (Medical Devices and Pharmaceutical Industry).
- To collaborate in projects on GxP compliance and Validation of computer systems, equipment, drug products & API processes and Quality Systems Development.
- Strong team-work skills are required since projects are carried out by interdisciplinary teams combining company employees and customer staff. Project Management experience is an added value.

Desired Skills And Experience

It is essential a Degree in Chemistry, Chemical Engineering, Biotechnology or Pharmacy.

Valuable two years working in pharmaceutical organizations under GxP compliance or similar.

The ideal candidate should have experience in GxP compliance, validation, quality system organization, medical device regulation.

Good level of knowledge of pharma and chemical business & manufacturing processes

Candidates should be very fluent in spoken and written English

The activities performed by TDV staff are mainly based at customers' sites, therefore the availability of a car is a must for the job.

Customers are mostly located within Spain, but also at European countries, China or other. Willingness to travel is mandatory.

We Offer

Stable position and a full time job

Projection within the company

Competitive salary

Contact

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