

### Medical and regulatory affairs associated

We are **Sycal Medical**! A young digital health start-up focused on increasing the detection of early-stage cancer and improving patient's life quality applying AI-based algorithms to medical imaging tests. Our first product helps radiologists to detect and classify pre-cancerous lesions in the abdomen on Medical Imaging tests.

Sycal is engaged in an active phase of development and hiring young talent to help us grow together! If this sounds like an exciting opportunity, we want to hear from you!

We are currently looking for a focused, autonomous, and ambitious PhD in Health Science who wishes to pursue a career in the industrial sector. In this position you will work on developing **clinical studies**; scientific investigation, document preparation, information management, file maintenance, and coordination of tasks across multiple departments. Part of your strategic directive will be to achieve a balance between regulatory concerns, technology, marketing objectives, compliance, time to market, and costs. For this reason, communication with employees at all levels within the organization is extremely important!

As a Medical and regulatory affairs associated you will need to understand aspects of product development, including **research, clinical trials, regulations, and approval processes**. Your research developed at Sycal will bring healthcare professionals diagnostic improvements, treatment enhancements and a greater range of integrated responses to clinical situations.

#### What you participate in:

- State-of-the-art analysis.
- Design, set-up, coordinate, and follow-up of Clinical Trials.
- Provide technical review of data or reports for statistical analysis.
- Provide support in regulatory affairs.
- Assistance with local Competent Authorities communication.
- Conformity Assessment – Medical Device EU Certification Procedure assistance.
- Evaluation of applicable laws and regulations to determine impact on company activities.
- Coordination of regulatory documentation activities.

#### Who you are:

- You have a Bachelor's, Master's Degree or PhD in the disciplines of life science, clinical science, public health or similar.
- Experience with regulatory procedures and clinical trials
- Experience in scientific investigation
- Indispensable fluent English and Spanish, spoken and written.
- Organized and resolute.
- Agile, flexible and able to adapt to changes.

**What do we offer:**

- Permanent contract with 6 months probation
- Young, open innovative and dynamic environment in the center of Barcelona.
- Flexible work-life balance, balancing working hours and home office.
- Growing together: we are an early-stage company with a multidisciplinary team.
- Equal employment opportunity: we proudly pursue a diverse workforce and we do not make any hiring or employment decisions that could be discriminatory in any way.