

IEC-Sites is currently seeking for a “**Medical Monitor Assistant**” able to manage images and reports performed in clinical trials at IEC sites (Barcelona & Madrid), to be part of the Medical Department.

As an **imaging CRO**, we provide full services in clinical trials: schedule patients imaging tests, perform radiology and nuclear medicine reports, images QC and response assessment follow up, send images to Corelab and query resolution.

This is an **exciting opportunity** for motivated professional to join one of the Spanish largest providers of medical imaging services solutions for clinical trials, serving pharmaceutical industry and CROs.

#### Job Requirements:

- **Lifesciences degree** or similar.
- Should be a **highly motivated and dynamic professional** with a positive and ambitious attitude to make things happen.
- Creative mind with ability for suggesting improvements and a solid **organizational and planning skills** in a fast-paced environment.
- A solid **interpersonal and communicative skills** for interacting with the staff and other colleagues
- Understanding of databases and data analysis procedures (collection, analysis, distribution etc.).
- Microsoft office program literacy.
- Previous experience in multi-center trials using radiological endpoints will be highly valued
- Previous experience in oncology clinical research will be positively valued
- Overall comprehensive knowledge of GCP and QC as it applies to clinical trials will be valued
- Oral and written fluency in Spanish and English.

#### Key Responsibilities:

- Knowing the general procedures of IEC-Sites studies in accordance with internal SOPs, ICH, GCP, relevant guidelines and all applicable laws and regulations.
- Reading radiological and nuclear medicine reports, for elaborating the QC and the patients' follow-up tables (RECIST, CHESON...).
- Respond medical queries related to images, medical reports, or follow-up tables.
- Supporting CTA-I for resolving doubts regarding the different imaging modalities, imaging reports and clinical data.



**Conditions:**

- Permanent Contract
- Full time position (40h / week).
- Career development.
- Immediate start