| JOB OFFER | | |
|---|---|---|
| **REFERENCE** | **OPENING DATE** | **DEADLINE** |
| 2018.246 | 24/09/2021 | 03/10/2021 |

| PROFILE REQUIREMENTS | |
|---|---|
| **EXCLUSIVE REQUIREMENTS: (1)** | |
| **ACADEMIC DEGREE** | Specialist in Clinical Pharmacology<br>Graduate in Medicine |
| **EXPERIENCE** | Support for clinical oncology trials in all their phases |
| **VALUED MERITS /SKYLLS** | |
| **EXPERIENCE** | Experience in database management<br>Experience in monitoring, and data manager in clinical trials, especially in trials in early phases<br>Experience in design and implementation of research projects<br>Experience in Integration in a multidisciplinary group of biomedical research<br>Experience in advise and contribute to the training of technical staff of the group |

| CONTRACT INFORMATION | | |
|---|---|---|
| **TYPE OF CONTRACT** | **EXPECTED INCORPORATION DATE** | **JOB STATUS** |
| A research project | octubre | Full time 40h/week |

| ANNUAL GROSS SALARY | DURATION OF THE CONTRACT |
|---|---|
| 46.089,91€ | 6 months (extendable depending on depending on the project and economic availability) |
| **WORK LOCATIONS** | **UNIT/DEPARTMENT** |
| Edif. Valdecilla Sur, 2ª planta, consultas de oncología | Servicio Oncología Médica HUMV |

| OFFER DESCRIPTION |
|---|
| Research support technician |

**DESCRIPTION OF THE TASKS IN THE PROJECT**

- Collection of data from medical records, obtaining informed consent
- Management of databases for statistical analysis
- Analysis of results
- Writing of manuscripts in English
- Assessment of the inclusion criteria of the patients for inclusion in the trial
- Toxicity assessment and dose adjustment
- Preparation of protocols for administration and management of the product under investigation
- Support in monitoring, pharmacovigilance and data collection. Resolution of doubts of the monitoring personne

| PRINCIPAL INVESTIGATOR / RESPONSABLE | RESEARCH GROUP | RESEARCH PROJECT |
|---|---|---|
| Fernando Rivera Herreo | Medical Oncology Service HUMV | 2020.407 Ensayo Fase III, Randomizado, Doble ciego y controlado con Placebo de Durvalumab y Quimioterapia FLOT Neoadyuvante-Adyuvante seguido de Durvalumab Adyuvante en pacientes con cáncer gástrico y de la unión gastroesofágica resecable (GC/GEJC) |

| | | |
|---|---|---|
| | | **(MATTERHORN)** |

| **RECRUITMENT INFORMATION** | |
|---|---|
| **SELECTION PROCESS STAGES (2)** | **EMPLOYMENT EXCHANGE** |
| **Pre-selection**<br>**Interview: maximum number of candidates to be interviewed: 3. Minimum score for this phase: 40.**<br>**Report of the Selection Board**<br>**Resolution** | **NOT** |

**SELECTION BOARD**

- **Galo Peralta, Director de Gestión de IDIVAL**
- **Patricia Álvarez, Coordinadora de Recursos Humanos (actuará de secretaria del Tribunal)**
- **Fernando rivera Herrero, Investigador Responsable del Proyecto**

**VALUATION OF MERITS**

| MERITS | EVALUATION | SCORE | | MAXIMUM |
|---|---|---|---|---|
| **Experience in database management** | **Curriculum** | **Compliance with the requirement** | **Yes/No** | **10** |
| **Experience in monitoring, and data manager in clinical trials, especially in trials in early phases** | **Curriculum** | **Compliance with the requirement** | **Yes/No** | **10** |
| **Experience in monitoring, and data manager in clinical trials, especially in trials in early phases** | **Curriculum** | **Compliance with the requirement** | **Yes/No** | **10** |
| **Experience in Integration in a multidisciplinary group of biomedical research** | **Curriculum** | **Compliance with the requirement** | **Yes/No** | **15** |
| **Experience in advise and contribute to the training of technical staff of the group** | **Curriculum** | **Compliance with the requirement** | **Yes/No** | **15** |

| **FINAL SCORE** | |
|---|---|
| **MAXIMUM TOTAL SCORE BY MERITS** | **60** |
| **MAXIMUM TOTAL SCORE IN INTERVIEW** | **40** |
| **MAXIMUM TOTAL SCORE** | **100** |

(1) Not subsanable
(2) See duration of each phase in the document "Selection Process"

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