

## JOB OFFER

REFERENCE	OPENING DATE	DEADLINE		
2015.054	12/09/2018	21/09/2018		
<b>WORKPLACE</b>				
<b>RESEARCH GROUP</b>	<b>PRINCIPAL INVESTIGATOR</b>			
Clinical Trials, Medical Oncology and Palliative Medicine	Fernando Rivera Herrero			
<b>WORKPLACE</b>	<b>UNIT / DEPARTMENT</b>			
HUMV	Servicio Oncología Médica HUMV			
LOCATION WORK PLACE (building, pavilion, plant etc.)	LOCALITY	POST CODE		
Edif. Valdecilla Sur, 2 <sup>a</sup> planta, consultas de oncología	Santander	39008		
<b>PROFILE REQUIREMENTS -2 POSITIONS</b>				
<b>PROFESSIONAL CATEGORY</b>	<b>ACADEMIC DEGREE</b>			
Medical Specialist	Bachelor of Medicine, specialist in Clinical Pharmacology			
<b>CANDIDATE REQUIREMENTS</b>				
<ul style="list-style-type: none"> <li>Extensive experience in supporting clinical trials in the area of clinical oncology in all its phases</li> </ul>				
<b>Valued merits / skills</b>				
<ul style="list-style-type: none"> <li>Medium level of English</li> <li>Experience in database management</li> <li>Experience in monitoring, and data manager in clinical trials, especially in trials in early phases</li> <li>Experience in design and implementation of research projects</li> <li>Integration capacity in a multidisciplinary group of biomedical research</li> <li>Ability to advise and contribute to the training of technical staff of the group</li> <li>Capacity for teamwork</li> </ul>				
<b>RECRUITMENT INFORMATION</b>				
<b>RESEARCH PROJECT</b>				
Project PCYC -1128-CA: Phase 1b / 2 study of combined treatment with ibrutinib in selected advanced gastrointestinal and genitourinary tumors.				
<b>DESCRIPTION OF THE TASKS IN THE PROJECT</b>				
<ul style="list-style-type: none"> <li>Collection of data from medical records, obtaining informed consent</li> <li>Management of databases for statistical analysis</li> <li>Analysis of results</li> <li>Writing of manuscripts in English</li> <li>Assessment of the inclusion criteria of the patients for inclusion in the trial</li> <li>Toxicity assessment and dose adjustment</li> <li>Preparation of protocols for administration and management of the product under investigation</li> <li>Support in monitoring, pharmacovigilance and data collection. Resolution of doubts of the monitoring personnel</li> </ul>				

- **Communication with the international team to resolve incidents**

DURATION OF THE CONTRACT	JOB STATUS	ANNUAL GROSS SALARY IN FULL TIME (40H PER WEEK)
<b>6 Months (provided funding exists) extendable for longer periods</b>	<b>40h/week</b>	<b>43.690,87€</b>

**SELECTION COURT**

- **Fernando Rivera , Project´s Main Researcher**
- **Galo Peralta, IDIVAL´s Management Director**
- **Marta Abelleira, Human Resources Coordinator (She will act as registrar of the selection board)**

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