

## JOB OFFER

REFERENCE	OPENING DATE	DEADLINE	
PT17/0017/0039	19/06/2019	28/06/2019	
<b>WORKPLACER</b>			
<b>RESEARCH GROUP</b>		<b>PRINCIPAL INVESTIGATOR</b>	
Central support Unit		María del Mar García Sáiz	
<b>WORKPLACE</b>		<b>UNIT/DEPARTMENT</b>	
HUMV		Clinical Trials Unit Valdecilla	
<b>LOCATION WORK PLACE (building, pavilion, plant etc.)</b>		<b>LOCALITY</b>	<b>Post Code</b>
Clinical Trials Unit Valdecilla. Pavilion 15-2 <sup>nd</sup> . HUMV		Santander	39008
<b>PROFILE REQUIREMENTS</b>			
<b>PROFESSIONAL CATEGORY</b>		<b>ACADEMIC DEGREE</b>	
Research Support Technician		Third Cycle Doctoral Degree	
<b>CANDIDATE REQUIREMENTS</b>			
<ul style="list-style-type: none"> <li>It is indispensable to have experience supporting and monitoring clinical trials. Minimal experience of 4 years.</li> <li>Good Clinical Practice (GCP) certification</li> <li>It is required to have at least an intermediate level of spoken and written English.</li> </ul>			
<b>VALUED MERITS/SKYLLES</b>			
<ul style="list-style-type: none"> <li>Formal training in Clinical Trials: Master</li> <li>Appropriate training and experience in monitoring, management and support to clinical trials and, in general, to clinical research from the point of view of management.</li> <li>Proven experience in management of medication and samples from patients, within clinical trials.</li> </ul>			
<b>RECRUITMENT INFORMATION</b>			
<b>RESEARCH PROJECT</b>			
PT17/0017/0039. SCReN- Platform for Clinical Research and Clinical Trial Units, project funded by Instituto de Salud Carlos III (the Ministry of Science and Innovation and Universities) and co-funded by FEDER.			
<b>DESCRIPTION OF THE TASKS IN THE PROJECT</b>			
<p>Support to IDIVAL Research Groups, according to the ISCIII SCREN platform rules, in the monitoring, management and implementation of clinical trials and in their follow-up. This includes supporting for the preparation of trial documentation, request for authorizations, monitoring, support in the reporting of adverse events/reactions and preparation of reports.</p> <p><b>It also includes collaborating with the Valdecilla Clinical Trials Unit in the preparation</b></p>			

**of proposals for clinical research projects and in the operational development of Clinical Trials.**

DURATION OF THE CONTRAT	JOB STATUS	ANNUAL GROSS SALARY IN FULLTIME
6 months extendable depending on the project and the economic availability	Full-time	26.395,17€

**SELECTION BOARD**

- **María del Mar García, Project´s Main Researcher**
- **Galo Peralta, IDIVAL´s Management Director**
- **Patricia Alvarez, Human Resources Coordinator.**

**A personal interview can be developed for the candidates with the best merit assessment.**

In compliance with the provisions of the Spanish Organic Law 15/1999 on Data Protection, of December 13, we inform you that the personal data provided to IDIVAL (hereinafter the Entity), will be included in an automated personal data filing system owned by the latter and kept under their responsibility, in order to manage their participation in our personnel selection processes. You may exercise the right of objection, access, rectification and erasure in relation to your personal data by writing to IDIVAL's Information Department through the email [idual@idual.org](mailto:idual@idual.org)