

<b>JOB OFFER</b>		
<b>REFERENCE</b>	<b>OPENING DATE</b>	<b>DEADLINE</b>
<b>PT20/00084</b>	<b>27/05/2021</b>	<b>05/06/2021</b>
<b>PROFILE REQUIREMENTS</b>		
<b>EXCLUSIVE REQUIREMENTS: (1)</b>		
<b>ACADEMIC DEGREE</b>	<b>Degree in Health Sciences + Official Master /equivalent</b>	
<b>EXPERIENCE</b>	<b>Experience in supporting the development of Clinical Trials.</b>	
<b>VALUED MERITS /SKYLLS</b>		
<b>FURTHER</b>	<b>Specialist in Clinical Pharmacology Good Clinical Practice Certification</b>	
<b>LANGUAGES</b>	<b>English (nivel ≥ B1)</b>	
<b>CONTRACT INFORMATION</b>		
<b>TYPE OF CONTRACT</b>	<b>EXPECTED INCORPORATION DATE</b>	<b>JOB STATUS</b>
<b>To research Project</b>	<b>June, 15</b>	<b>Full time (35h/week)</b>
<b>ANNUAL GROSS SALARY IN FULL TIME</b>	<b>DURATION OF THE CONTRACT</b>	
<b>22.247,36€</b>	<b>1 year (extendable depending on the project and financial availability)</b>	
<b>WORK LOCATIONS</b>	<b>UNIT/DEPARTMENT</b>	
<b>Hospital Universitario Marqués de Valdecilla. IDIVAL. Pavilion 15-2º</b>	<b>Area of Clinical Trials. Clinical Pharmacology Service</b>	
<b>OFFER DESCRIPTION</b>		
<b>Research Technician</b>		
<b>DESCRIPTION OF THE TASKS IN THE PROJECT</b>		
<ul style="list-style-type: none"> <li>- To establish a specific pharmacovigilance plan for the assigned clinical trials and to monitor the compliance of the plan and the Standard Operating Procedures.</li> <li>-To record and evaluate adverse event reports received from the investigators.</li> <li>-To enter the adverse event notifications in the specific data management and information system.</li> <li>-To perform expedite reporting of the suspected unexpected serious adverse reactions of clinical trials.</li> <li>-To prepare the clinical trial periodic safety reports.</li> <li>-To identify and evaluate relevant safety issues of investigational medicinal products or devices in clinical trials.</li> <li>- To collaborate with the design and the implementation of pharmacovigilance and pharmacoepidemiology research projects.</li> <li>- To assist the activities of the clinical research projects under the coordination of the Project Manager.</li> </ul>		
<b>PRINCIPAL INVESTIGATOR / RESPONSABLE</b>	<b>RESEARCH GROUP</b>	<b>RESEARCH PROJECT</b>
<b>Mª del Mar García Sáiz</b>	<b>Area of Clinical Trials. Clinical Pharmacology Service</b>	<b>Spanish Clinical Research Network (SCReN) "Aid financed by the Instituto de Salud Carlos III (Ministry of Science and Innovation) and co-financed by the European Regional Development Fund (ERDF)".</b>

RECRUITMENT INFORMATION				
SELECTION PROCESS STAGES (2)				EMPLOYMENT EXCHANGE
<b>Preselection:</b> <b>Inerview: maximum number of candidates to be interviewed: 4. Minimum score for this stage: 10</b> <b>Tribunal report:</b> <b>Resolution:</b>				NOT
SELECTION BOARD				
<ul style="list-style-type: none"> <li>• M<sup>a</sup> del Mar García Sáiz, Project´s Main Researcher</li> <li>• Galo Peralta, IDIVAL´s Management Director</li> <li>• Patricia Álvarez-Ingelmo, IDIVAL Human Resources Coordinator (She will act as registrar of the selection board).</li> </ul>				
VALUATION OF MERITS				
MERITS	EVALUATION	SCORE		MAXIMUM
Specialist in Clinical Pharmacology	Supporting document	Requirement fulfilment	Yes/No	40
Good Clinical Practice Certification	Supporting document	Requirement fulfilment	Yes/No	15
English	Supporting document	Level ≥B1	Yes/No	5
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"

*In compliance with the provisions of Article 11 of Organic Law 3/2018, you are informed that the person responsible for the processing of your personal data is the MARQUES DE VALDECILLA INSTITUTE OF INVESTIGATION FOUNDATION (IDIVAL), your data will be treated in order to be treated to the extent that they were necessary or convenient for the development of the legal relationship established between the parties. You can exercise your rights of access, rectification, deletion, opposition, portability or limitation of the treatment, by contacting the IDIVAL FOUNDATION at the following address: AVDA. CARDENAL HERRERA ORIA, S / N 39007, SANTANDER. More information at [www.idival.org/es/Política-de-Privacidad](http://www.idival.org/es/Política-de-Privacidad)*