

JOB OFFER		
REFERENCE	OPENING DATE	DEADLINE
<b>ICI25/0088</b>	<b>03/03/2026</b>	<b>12/03/2026</b>
PROFILE REQUIREMENTS		
EXCLUSIVE REQUIREMENTS: (1)		
ACADEMIC DEGREE	<b>Bachelor's degree + Official Master's Degree/equivalent (Proof must be provided with the application).</b>	
CONTRACT INFORMATION		
TYPE OF CONTRACT	EXPECTED INCORPORATION DATE	JOB STATUS
<b>Contract for scientific-technical activities</b> (article 23.bis of Law 14/2011, of June 1, on Science, Technology and Innovation)	<b>March 2026</b>	<b>Full time.</b> 1575 hours per year (aprox. 35h/week)
ANNUAL GROSS SALARY	DURATION OF THE CONTRACT	
<b>28.755,20€</b> , without prejudice to any adjustment that may be established in state legislation for 2026.	<b>Indefinite</b> (linked to the duration of the project or to external financing or financing from public grants in full competition).	
WORK LOCATIONS	UNIT/DEPARTMENT	
<b>HUMV Pavilion 15-2nd Floor</b>	<b>Clinical Trials Area. Clinical Pharmacology Service</b>	
JOB DETAILS		
OFFER DESCRIPTION		
<b>Research support technician</b>		
FUNCTIONS		
<ul style="list-style-type: none"> <li>-Ensure compliance with Good Clinical Practice standards, applicable legislation and Standard Operating Procedures.</li> <li>-Ensure effective communication between the research team/sponsor at the assigned centres.</li> <li>-Monitor the clinical research project activities carried out at the assigned centres as defined by the project manager.</li> <li>-Support the Project Manager and CRA in those activities required for the proper conduct of the clinical trial.</li> <li>-Prepare periodic review reports and updates for the project manager/sponsor.</li> <li>-Keep essential project documentation up to date and properly filed at the centres assigned to you.</li> <li>-Perform data analysis according to project needs.</li> <li>-Construction of data collection notebooks for clinical trials.</li> <li>-Obtain clinical data or records from project management tools under the coordination of the project manager.</li> <li>-Support for resolving inconsistencies, deviations and errors in the data collected from the clinical trial (queries).</li> <li>-Ensure the traceability of data at the centres assigned to you.</li> <li>-Actively collaborate in ensuring the quality of data, documentation and processes at the assigned centres.</li> </ul>		
PRINCIPAL INVESTIGATOR / RESPONSABLE	RESEARCH GROUP	RESEARCH PROJECT
<b>Dr. Enrique M. Ocio San Miguel</b>	<b>Haematological malignancies and haematopoietic stem cell transplantation</b>	<b>ICI25/0088. Safety and efficacy of allogeneic a-iNKT cell adoptive immunotherapy in elderly patients with intermediate/adverse-risk AML who have had a suboptimal response to venetoclax and azacitidine.</b>
RECRUITMENT INFORMATION		
SELECTION PROCESS STAGES (2)		EMPLOYMENT EXCHANGE

<b>1. Admission of applications.</b> <b>2. Competition phase.</b> <b>3. Interview phase:</b> maximum number of candidates to be interviewed: 4. Minimum score for this phase:40 <b>4. Report of the Tribunal.</b> <b>5. Resolution.</b>				<b>NOT</b>
<b>NOTE: in order for candidates to be considered for recruitment and employment exchange purposes, they must have a total score of at least 30 points.</b>				
<b>SELECTION BOARD</b>				
<ul style="list-style-type: none"> <li>• <b>President:</b> Enrique M. Ocío San Miguel, Principal Investigator of the Project</li> <li>• <b>Member:</b> M. Blanca Sanchez Santiago – Acting Head of Clinical Pharmacology</li> <li>• <b>Member and Secretary:</b> Lucía Lavín Alconero, Clinical Trials Assistant Technician.</li> </ul>				
<b>VALUATION OF MERITS</b>				
MERITS	EVALUATION	SCORE		MAXIMUM
Training in data management or data science	Supporting document	Merit compliance	YES/NO	25
Experience in database design	Curriculum vitae	Merit compliance	YES/NO	5
Experience in REDCAP design	Curriculum vitae	Merit compliance	YES/NO	10
Data analysis (statistics)	Curriculum vitae	Merit compliance	YES/NO	10
Certification in Good Clinical Practice R3	Supporting document	Merit compliance	YES/NO	5
Scientific publications	Curriculum vitae	Merit compliance	YES/NO	5
<b>FINAL SCORE</b>				
<b>MAXIMUM TOTAL SCORE BY MERITS</b>				<b>60</b>
<b>MAXIMUM TOTAL SCORE IN INTERVIEW</b>				<b>40</b>
<b>MAXIMUM TOTAL SCORE</b>				<b>100</b>

- (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/Politica-de-Privacidad](http://www.idival.org/es/Politica-de-Privacidad)*

Santander as of the date of electronic signature

Fdo. Francisco Galo Peralta Fernandez