

JOB OFFER		
REFERENCE	OPENING DATE	DEADLINE
EU25/05	07/07/2025	16/07/2025
PROFILE REQUIREMENTS		
EXCLUSIVE REQUIREMENTS: (1)		
ACADEMIC DEGREE	Bachelor's degree + Official Master's degree/equivalent (must be justified with the application).	
CONTRACT INFORMATION		
TYPE OF CONTRACT	EXPECTED INCORPORATION DATE	JOB STATUS
Contract for scientific-technical activities (article 23.bis of Law 14/2011, of June 1, on Science, Technology and Innovation)	July 2025	Full time. 1575 hours per year (aprox. 35h/week)
ANNUAL GROSS SALARY	DURATION OF THE CONTRACT	
26.717,29€, without prejudice to the basic update established in state legislation for 2024.	Indefinite (linked to the duration of the project or to external financing or financing from public grants in full competition).	
WORK LOCATIONS	UNIT/DEPARTMENT	
Hospital Universitario Marqués de Valdecilla. Pabellón 15-2ª planta	Clinical Trials Area. Clinical Pharmacology Service	
JOB DETAILS		
OFFER DESCRIPTION		
Research support technician		
FUNCTIONS		
<ul style="list-style-type: none"> <li>- To monitor the clinical trial activities carried out at the assigned sites as defined by the project manager.</li> <li>- Ensure effective communication between the investigator/promoter team at the assigned sites.</li> <li>- Conduct the kick-off visit and train the research team in the project activities.</li> <li>- To carry out the monitoring in compliance with the Monitoring Plan and manual,</li> <li>- Verify that the protocol and its modifications are complied with.</li> <li>- Ensure compliance with Good Clinical Practice guidelines, applicable legislation and Standard Operating Procedures.</li> <li>- Carrying out the clinical trial closure visits.</li> <li>- Prepare monitoring and review reports to the project manager.</li> <li>- Keep essential trial documentation up to date and correctly filed at assigned sites.</li> <li>- Support for the resolution of inconsistencies, deviations and errors in the data collected from the trial (queries).</li> <li>- Ensure traceability of medication delivered to assigned sites.</li> <li>- Maintain the information required from the assigned sites in the project monitoring tools.</li> <li>- Actively collaborate in the quality assurance of the data, documentation and processes of the centres.</li> <li>- Produce periodic reports in English as defined by the project manager.</li> <li>- Collaboration in product materiovigilance reports.</li> </ul>		
PRINCIPAL INVESTIGATOR / RESPONSABLE	RESEARCH GROUP	RESEARCH PROJECT
María del Mar García Saiz	UEECC	EU25/05 MORPHEUS/ETHER: Prognosis improvement of unprovoked venous ThromboEmbolicism with the use of a shared decision-making process including a time-dependent multicomponent risk prediction scores intervention.
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: 3. Minimum score for this phase: 20 <b>4. Report of the Tribunal.</b> <b>5. Resolution.</b>  <b>Note:</b> 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>YES</b>
<b>SELECTION BOARD</b>				
<ul style="list-style-type: none"> <li>• <b>President:</b> María Blanca Sánchez Santiago, Head of the Clinical Trials Unit..</li> <li>• <b>Member:</b> Lucía Lavín Alconero, project partner</li> <li>• <b>Member and secretary:</b> : Maria José Marín Vidalled, IDIVAL Technology Services Coordinator</li> </ul>				
<b>VALUATION OF MERITS</b>				
<b>MERITS</b>	<b>EVALUATION</b>	<b>SCORE</b>		<b>MAXIMUM</b>
Master in Clinical Trials/ Master in Health Care Research	Documentary evidence	Compliance with the merit	Yes/No	<b>15</b>
Good Clinical Practice Certification	Documentary evidence	Compliance with the merit	Yes/No	<b>10</b>
Experience in monitoring clinical trials / Experience in pharmacovigilance/materiovigilance management of clinical trials.	Curricular	Compliance with the merit	Yes/No	<b>15</b>
Experience as Data Manager or Study Coordinator in Clinical Trials	Curricular	Compliance with the merit	Yes/No	<b>10</b>
English B2 or higher	Documentary evidence	Compliance with the merit	Yes/No	<b>5</b>
Driving licence B	Curricular	Compliance with the merit	Yes/No	<b>5</b>
<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"

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Santander as of the date of electronic signature

Fdo. Francisco Galo Peralta Fernandez

