

JOB OFFER		
REFERENCE	OPENING DATE	DEADLINE
2016.188	01/04/2026	10/04/2026
PROFILE REQUIREMENTS		
EXCLUSIVE REQUIREMENTS: (1)		
ACADEMIC DEGREE	PhD in Health Sciences or equivalent (Proof must be provided with the application).	
EXPERIENCE	Experience in translational research: observational studies and clinical trials	
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)	MAYO 2026	Part time. 1575 hours per year (aprox. 35h/week)
ANNUAL GROSS SALARY	DURATION OF THE CONTRACT	
33.139,97€ without prejudice to the basic update established in state legislation for 2026.	Indefinite (linked to the duration of the project or to external financing or financing from public grants in full competition).	
WORK LOCATIONS	UNIT/DEPARTMENT	
HUMV	Infectious Diseases	
JOB DETAILS		
OFFER DESCRIPTION		
Research support technician		
FUNCTIONS		
<ul style="list-style-type: none"> - Completing data collection forms. - Managing incoming and outgoing information. - Coordinating clinical research within the department: observational studies and clinical trials. - Liaising with clinical trial monitors and sponsors; and with other departments involved in the projects. - Processing and dispatching biological samples. - Maintaining and updating study documentation. - Supporting the infectious disease specialist in the conduct of the clinical trial as Study Coordinator and Data Manager (patient selection, scheduling visits, completing CRFs, resolving queries). - Assisting with coordination tasks in multicentre trials. - Assisting in the preparation of project applications, interim reports and final reports. - Assisting in the preparation of reports on the group's research activities. - Designing and producing data collection forms for studies. - Assisting in data analysis and the drafting of manuscripts. 		
PRINCIPAL INVESTIGATOR / RESPONSABLE	RESEARCH GROUP	RESEARCH PROJECT
María Carmen Fariñas Álvarez	Infectious Diseases and Clinical Microbiology	2016.188 A Phase III, randomised, multicentre, open-label, parallel-group study to evaluate the efficacy, safety and tolerability of a long-acting intramuscular regimen containing cabotegravir and rilpivirine in maintaining virological suppression, following induction with a single-tablet regimen containing an integrase inhibitor in a Phase III, randomised, multicentre, open-label, parallel-group study to evaluate the efficacy, safety and tolerability of a long-acting intramuscular regimen of cabotegravir and rilpivirine in maintaining virological suppression, following induction with a single-tablet regimen containing an integrase inhibitor in HIV-1-infected adult patients who have not received prior antiretroviral therapy HIV-1-infected adult patients who have not received prior antiretroviral therapy



RECRUITMENT INFORMATION				
SELECTION PROCESS STAGES (2)			EMPLOYMENT EXCHANGE	
1. Admission of applications. 2. Competition phase. 3. Interview phase: maximum number of candidates to be interviewed:5. Minimum score for this phase:40 4. Report of the Tribunal. 5. Resolution.			NOT	
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.				
SELECTION BOARD				
<ul style="list-style-type: none"> • President: Dra. María Carmen Fariñas Álvarez Assistant Researcher. • Member: Marcos López Hoyos, Scientific Director • Member and secretary: María José Marín, IDIVAL SSTT Coordinator. 				
VALUATION OF MERITS				
MERITS	EVALUATION	SCORE		MAXIMUM
Involvement in grants, contracts and research projects in the field of biomedicine*	Curriculum (please specify duration)	Achievement of merit	Up to 5 years: 4 points Between 5 and 10 years: 8 points Over 10 years: 12 points	12
Experience in project management*	Curriculum (please specify duration)	Achievement of merit	Up to 5 years: 4 points Between 5 and 10 years: 8 points More than 10 years: 12 points	12
Proficiency in IT tools (REDCap, Access) or basic knowledge of bioinformatics	Curriculum	Achievement of merit	Yes/No	10
Certificate in Good Clinical Practice	Curriculum	Achievement of merit	Yes/No	3
Knowledge of statistics	Curriculum	Achievement of merit	Yes/No	12
Experience in coordinating observational studies and clinical trials	Curriculum	Achievement of merit	Yes/No	6
Experience in data collection for observational studies and clinical trials and responding to queries	Curriculum	Achievement of merit	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"

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Santander as of the date of electronic signature

Fdo. Francisco Galo Peralta Fernandez

Av. Cardenal Herrera Oria s/n | www.idival.org
 39011 Santander - España | Tel. +34 942 31 55 15

Fundación Instituto de Investigación Marqués de Valdecilla
 CIF: G 39788773

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FIRMANTE(1) : FRANCISCO GALO PERALTA FERNÁNDEZ | FECHA : 30/03/2026 12:07 | Sin acción específica

