

JOB OFFER		
REFERENCE	OPENING DATE	DEADLINE
2020.279	01/06/2023	10/06/2023
PROFILE REQUIREMENTS		
EXCLUSIVE REQUIREMENTS: (1)		
ACADEMIC DEGREE	Medical Specialist in Clinical Pharmacology (Excluding requirement: justification must be provided with the application).	
OTHERS REQUIREMENTS	B2 level of English (Excluding requirement: justification must be provided with the application).	
VALUED MERITS /SKYLLS		
EXPERIENCE	<ul style="list-style-type: none"> - Experience as clinical trial coordinator - Experience as an investigator-subinvestigator in clinical trials 	
OTHERS	<ul style="list-style-type: none"> - Training in the field of clinical trials (good clinical practices in clinical research, monitoring of clinical trials, IATA, etc.). - Training in Emergency Medicine - Research (publications and conference communications) in the field of clinical pharmacology 	
CONTRACT INFORMATION		
TYPE OF CONTRACT	EXPECTED INCORPORATION DATE	JOB STATUS
Indefinite in accordance with Article 23 Bis of Law 14/2011, of June 1, 2011, on Science, Technology and Innovation	June	Full-time (35h/week)
ANNUAL GROSS SALARY IN FULL TIME		DURATION OF THE CONTRACT
42.783,68 €		Indefinite term linked to the duration of the project and to the economic availability based on RDL8/2022.
WORK LOCATIONS		UNIT/DEPARTMENT
IDIVAL		Clinical trials unit Valdecilla
OFFER DESCRIPTION		
Research support technician		
DESCRIPTION OF THE TASKS IN THE PROJECT		
<p>- Support the development of clinical trials throughout their development with the following tasks:</p> <ul style="list-style-type: none"> • Coordination and comprehensive planning of patients' agendas for their participation in clinical trials. • Attention to trial subjects in the Valdecilla Clinical Trials Unit, and specifically assistance in emergency and urgent situations. • Monitoring of the quality system • Interlocution with the investigators and monitors for the development of clinical trials. • Supervision of the tasks of drug administration, test performance, sample shipment, etc. o Supervision of the information systems of the clinical trials. • Supervision of the information systems used in the clinical trials. 		

PRINCIPAL INVESTIGATOR / RESPONSABLE		RESEARCH GROUP		RESEARCH PROJECT	
M ^a Blanca Sánchez Santiago		Clinical trials Unit Valdecilla		2020.279: Estudio fase 2a, aleatorizado, doble ciego, controlado con placebo, para evaluar un rango de niveles de dosis e intervalos de vacunación de Ad26CCOV51 en adultos sanos de 18 a 55 años inclusive	
RECRUITMENT INFORMATION					
SELECTION PROCESS STAGES (2)					EMPLOYMENT EXCHANGE
Pre-selection Interview: maximum candidates to interview: 3. Minimum score for this phase: 40 Report of the Selection Board Resolution					NOT
SELECTION BOARD					
<ul style="list-style-type: none"> • Blanca Sánchez Santiago, Project´s Main Researcher • Galo Peralta, IDIVAL´s Management Director • Patricia Álvarez-Ingelmo, IDIVAL Human Resources Coordinator (She will act as registrar of the selection board). 					
VALUATION OF MERITS					
MERITS	EVALUATION	SCORE		MAXIMUM	
Experience as clinical trial coordinator	Curricular	Merit fulfillment	YES/NO	20	
Experience as an investigator-subinvestigator in clinical trials	Curricular	Merit fulfillment	YES/NO	10	
Training in the field of clinical trials (good clinical practices in clinical research, monitoring of clinical trials, IATA, etc.).	Curricular	Merit fulfillment	YES/NO	10	
Training in Emergency Medicine	Curricular	Merit fulfillment	YES/NO	10	
Research (publications and conference communications) in the field of clinical pharmacology	Curricular	Merit fulfillment	YES/NO	10	
FINAL SCORE					
MAXIMUM TOTAL SCORE BY MERITS					60
MAXIMUM TOTAL SCORE IN INTERVIEW					40
MAXIMUM TOTAL SCORE					100

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- (1) Not subsanable
 - (2) See duration of each phase in the document "Selection Process"

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