JOB OFFER		
REFERENCE	**OPENING DATE**	**DEADLINE**
2017.061	**18/05/2023**	**27/05/2023**

PROFILE REQUIREMENTS	
EXCLUSIVE REQUIREMENTS: (1)	
ACADEMIC DEGREE	**Degree in health sciences *(Excluding requirement: provide justification with the application)***
VALUED MERITS /SKYLLS	
TRAINING	• **Office automation advanced user level.**
EXPERIENCE	• **Experience and knowledge in clinical trials.** • **Experience in CRDe management and queries resolution.** • **Database maintenance.**
LANGUAGES	• **English level.**

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	DURATION OF THE CONTRACT
21.543,68 €	**It will depend on the duration of the project and the economic availability based on RDL8/2022.**
WORK LOCATIONS	**UNIT/DEPARTMENT**
Pavilion 20, Hematology consultation rooms and floor. Clinical Trials Unit	**Hematology/Clinical Trials**

JOB DETAILS
OFFER DESCRIPTION
Research Support Technician
NUMBER OF POSITIONS OFFERED
2
DESCRIPTION OF THE TASKS IN THE PROJECT
• **Knowledge of the guidelines of the different notebooks and compliance with data entry metrics schedule.** • **Completion of specific training related to the position.** • **Knowledge of the protocols of the different clinical trials of the service, test schedules and their procedures.** • **Prepare monitoring visits and ensure proper maintenance of patient files.** • **Inventory laboratory materials provided by the sponsors and ensure their availability.** • **Patient education and monitoring in the use of technology (tablet, phone...) within the scheduled hospital visits.** • **Provide support to the coordinators of the different studies in the different daily procedures (sending samples, consents...).**

- **Other functions that may be determined.**

PRINCIPAL INVESTIGATOR / RESPONSABLE	RESEARCH GROUP	RESEARCH PROJECT
Dr Enrique María Ocio San Miguel	**Hematologic Neoplasms and Hematopoietic Progenitor Transplantation**	**2017.061: Estudio de fase III, doble ciego, controlado con placebo de quizartinib (AC220) administrado en combinación con quimioterapia de inducción y consolidación, y administrado como terapia de mantenimiento a sujetos de 18 a 75 años con leucemia mielógena aguda FLT3-ITD (+) de nuevo diagnóstico (QuANTUM First).**

RECRUITMENT INFORMATION	
SELECTION PROCESS STAGES (2)	**EMPLOYMENT EXCHANGE**
1. Preselection **2. Interview: maximum number of candidates to be interviewed: 5. Minimum score in merits to pass to the interview phase: 20.** **3. Report of the Truibunal** **4. Resolution**	**YES**

SELECTION BOARD

- **Enrique Ocio San Miguel, Principal Investigator of the Research Project**
- **Galo Peralta, Management Director of IDIVAL**
- **Patricia Álvarez-Ingelmo, HR coordinator (will act as Registrar of the Tribunal)**

VALUATION OF MERITS

MERITS	EVALUATION	SCORE		MAXIMUM
Office automation advanced user level	**Curriculum**	**Merit fulfillment**	**Yes / No**	**10**
Experience and knowledge in clinical trials	**Curriculum**	**Merit fulfillment**	**Yes / No**	**20**
Database maintenance	**Curriculum**	**Merit fulfillment**	**Yes / No**	**10**
Experience in CRDe management and queries resolution	**Curriculum**	**Merit fulfillment**	**Yes / No**	**10**
English	**Curriculum**	**Level**	**B1: 2 B2: 5 C1: 10**	**10**

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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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.