







	OPENIN 15/03/2022	IG DATE	DEADLINE		
	15/03/2022				
			24/03/2022		
	PROFILE REC	QUIREMENTS			
Е	XCLUSIVE REQ	UIREMENTS: (1	.)		
ACADEMIC DEGREE Bachelor's deg			ree + official master's degree/equivalent		
PERIENCE Research managen			jement experience		
OTHERS REQUIREMENTS English B1					
	VALUED MER	ITS /SKYLLS			
Training in monitoring, management, coordination and support of clinical trials and clinical research in general.					
 Management of international projects Management of consortium participation Experience in the processing of administrative documentation for ethics committees and clinical research in general. 					
English level higher than B1 and/or level in other languages other than English.					
 Training abroad Good Clinical Practice (GCP) accreditation. 					
	CONTRACT IN	NFORMATION			
TYPE OF CONTRACT			JOB STATUS		
Eventual APRIL 2022			Full time (35h/week)		
ANNUAL GROSS SALARY			DURATION OF THE CONTRACT		
24.940,88 €		6 months (extendable depending on the project and economic availability)			
WORK LOCATIONS		UNIT/DEPARTMENT			
IDIVAL/HUMV		CENTRAL SUPPORT UNIT			
	ining inical tria Man Man Expense for e	Bachelor's deg Research man NTS English B1 VALUED MER ining in monitoring, relical trials and clinical in Management of inte Management of core Experience in the properties committed in the	Bachelor's degree + official names of the Research management experience in monitoring, management, control trials and clinical research in gent of the Management of international project of Management of consortium particular of the Processing of a for ethics committees and clinical glish level higher than B1 and/or leven English. Training abroad Good Clinical Practice (GCP) accres the Contract Information of the Processing of the Project and Contract of the Project of the Proje		

OFFER DESCRIPTION

Research Support Technician

DESCRIPTION OF THE TASKS IN THE PROJECT

Its main functions and activities will be to support the coordination and management of the MIRCAST Project in collaboration with the principal investigator and the IDIVAL management team.

Activities will include:

- Management of legal documentation, agreements, contracts, etc.
- Administrative management.
- Follow-up of the progress of the project with the interlocution of the









researchers of the different nodes.

- Advising the researchers in monitoring the study.
- Preparation of the necessary follow-up documentation (technical reports, budgets, etc.).
- Preparation of justification documentation and, if necessary, audits.
- Dissemination of project highlights.
- Support in all those tasks related to the coordination of the MIRCAST project.

PRINCIPAL INVESTIGATOR / RESPONSABLE	RESEARCH GROUP	RESEARCH PROJECT	
Marcos Gómez Ruiz	Surgical Innovation Group	This plaza is convened within the framework of the multi-center research project "MIRCAST" of international scope whose promoter is IDIVAL and has public-private collaboration, whose initial planned duration is 3 years.	

RECRUITMENT INFORMATION

SELECTION PROCESS STAGES (2)	EMPLOYMENT EXCHANGE
Preselección Entrevista: máximo candidatos a entrevistar 4. Aquellos con la puntuación máxima. Informe del Tribunal Resolución	NOT

SELECTION BOARD

- Marcos Gomez Ruiz, Investigador Principal del Proyecto de Investigación
- Galo Peralta, Director de Gestión de IDIVAL
- Patricia Álvarez-Ingelmo, coordinadora de RRHH (actuará como Secretario del Tribunal)

VALUATION OF MERITS					
MERITS	EVALUATION	SCORE		MAXIMUM	
Training in monitoring, management, coordination and support of clinical trials and clinical research in general.	CV	Requirement fulfilment	YES/NOT	10	
International project management	CV	Requirement fulfilment	YES/NOT	10	
Management of consortium participation	CV	Requirement fulfilment	YES/NOT	10	
Experience in the processing	CV	Requirement fulfilment	YES/NOT	10	









of administrative documentation for ethics committees and clinical research in general.				
Training abroad	CV	Time	2 points for month	5
Level of English higher than B1 and/or level in languages other than English (LEVEL AND LANGUAGE)	Supporting document	Level	B2: 5 points C2 or superior: 10 points	10
Have accreditation in Good Clinical Practices (GCP).	Supporting document	Requirement fulfilment	YES/NOT	5
FINAL SCORE				
MAXIMUM TOTAL SCORE BY MERITS			60	
MAXIMUM TOTAL SCORE IN INTERVIEW			40	
MAXIMUM TOTAL SCORE			100	

⁽¹⁾ Not subsanable

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⁽²⁾ See duration of each phase in the document "Selection Process"