Parte A en las propuestas de HE

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Estructura de la Propuesta

- **Parte A:**
  - Formularios generados por el sistema, se rellenan en línea (Funding & Tender Portal).
  - Información general de la propuesta y los participantes.

- **Parte B:**
  - Parte narrativa. Descripción técnica de la propuesta.
  - Incluye 3 secciones que corresponden cada una a un criterio de evaluación.
  - Debe cargarse en el sistema como un documento PDF.

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Plantillas propuesta completa

Horizon Europe Programme

Standard Application Form (RIA, IA)

Application form (Part A)
Project proposal – Technical description (Part B)

Horizon Europe Programme

Standard Application Form (CSA)

Application form (Part A)
Project proposal – Technical description (Part B)

➢ Nuevas versión actualizada de las plantillas (*Version 4.0, 21 January 2022*)
➢ La plantilla se descarga del sistema de envío, específico para la convocatoria o *topic*.

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Roles en la propuesta

➢ **Coordinator**: actúa como único punto de contacto entre los socios del consorcio y la UE. El coordinador de la propuesta es, por lo general, la organización responsable de la planificación general de la propuesta; de la creación del consorcio y realizará el trabajo de administración de la propuesta en el sistema de presentación.

➢ **Partner (or participant)**: será invitado por el coordinador. Los contactos de los socios a los que se les conceden plenos derechos de acceso pueden rellenar los formularios administrativos relacionados con su entidad.

Roles en la propuesta

➢ **Affiliated Entities**: Los socios también pueden tener entidades afiliadas a ellos (Antiguas "Linked Third Parties" en H2020). Son entidades con un vínculo legal con los beneficiarios que ejecutan partes de la acción y pueden cargar los costes a la subvención. A nivel de la propuesta, no hay que proporcionar contactos para las entidades afiliadas y no pueden realizar ninguna acción administrativa en la propuesta electrónica.

➢ **Associated Partners**: Son entidades que ejecutan tareas de acción pero sin recibir financiación de la UE. A nivel de la propuesta, no hay que proporcionar sus contactos y no pueden realizar ninguna acción administrativa en la propuesta electrónica.

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Derechos de acceso a la propuesta

Role rights

Differences between a Proposal Coordinator and a Participant actions:

Action	Proposal Coordinator	Partner
Select the call	Yes	No
Add, Invite Participants	Yes	No
Submit the proposal	Yes	No
Read complete proposal	Yes	Yes
Define the budget tables	Yes	Yes (if full access has been selected for the contact), with rights to fill in only the row for their organisation.
Create Contacts for a Partner	Yes	Yes (if full access has been selected for the contact), with rights to fill in only the information for their organisation.

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➢ Todas las entidades que vayan a participar en la propuesta tienen que estar previamente registradas en el *Participant Register* a través del Funding & Tender Portal para obtener el PIC (*Participant Identification Code*).

Participant Register User´s Guide

https://ec.europa.eu/research/participants/data/support/manual/urf.pdf

Crear la propuesta en el F&T Portal

Funding & Tenders Portal (F&T Portal):
https://ec.europa.eu/info/funding-tenders/opportunities/portal/screen/home

➢ Proposal Submission Service User Manual
https://ec.europa.eu/research/participants/data/support/sep_usermanual.pdf

Terms and Conditions for the Submission of Applications ✕

Please read and accept this disclaimer to proceed with the creation of your draft proposal:

Proposal data

Confidentiality: Applications (and any additional information provided for the award) will be considered confidential and treated according to the same principles as those set out in the grant agreement. We may in particular have to share applications with other EU institutions and bodies in order to perform the pre-award checks required by the EU Financial Regulation. Please also be aware that we may have to publish the information contained in the Abstract/Short Summary in Part A. Please avoid adding any confidential information in that section.

Access to pre-registration data: Please be aware that certain information from the draft proposals will be accessed by us before the call deadline, in order to plan the evaluation. This concerns call ID, topic ID, type of action ID, participant identification code (PIC), project acronym, abstract, panel and keywords.

- ● I agree that the pre-registration data becomes available to the European Commission services prior to call closure.
- ○ I wish that the pre-registration data does not become available to the European Commission services prior to call closure. (Applicants are advised to use this option only in well justified cases, as it hinders the planning of the evaluation process and the timely processing of proposals).

Technical requirements

Part B file format: Part B of the Application Form (Technical Description) must be uploaded as PDF (equivalent to Adobe Acrobat v5 or higher, with embedded fonts).

Part B page limits: Applications are subject to the page limits set out in the call conditions. Excess pages will be made invisible and not available to the evaluators. Mandatory annexes and supporting documents do not count towards the page limit.

Time constraints: Preparation and uploading of the files in the Submission System may take some time. Don't wait until the end. Problems due to last minute submissions (e.g. congestion, etc) will be entirely at your risk.

Submission

Applications must be submitted before the call deadline. You can withdraw your application or make changes to it, provided this is done before the call deadline. Submission of a new application overwrites the previous version.

Personal Data

Personal data linked to the application, evaluation or award of grants, prizes and contracts will be processed in accordance with the Funding & Tenders Portal Terms and Conditions and Privacy Statement

Cancel   **OK**

Draft proposal TEST created ✕

Dear ▓▓▓▓▓▓
You have successfully created a draft proposal **TEST** for the call **ERASMUS-2021-EMJM-MOB**.

You can continue editing your draft proposal now or access it at a later time, but before the deadline of **31 January 2021 10:35:00 (Brussels Local Time)** from the Funding & Tenders Portal by accessing the My Proposals tab.

An email containing this information has been sent to this email address: ▓▓▓▓▓▓ (which is associated with your ▓▓▓ account ▓▓▓).

Go to My Proposals 🗗   **Continue with this proposal**

Una vez el borrador ha sido guardado, a la lista de "My Proposals" del Portal.
➢ "Go to My proposals" para volver al Portal
➢ "Continue with this proposal" para continuar con el siguiente paso

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Añadir participantes

> ➤ Si en el consorcio hay "Affiliated Entities" o "Associated Partners", se añadirán en el socio al que estén vinculados.

Añadir participantes

> Para volver a acceder a la propuesta desde la pantalla del Portal, ir al botón "Actions" de la propuesta y seleccionar "Edit Draft"

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Editar el formulario (Parte A de la propuesta)

< Step 5   Table of contents   General Information >

Application forms   **Table Of Contents**   Save   Save&Close

Promotion of ▮▮▮▮▮▮▮▮▮▮

▮▮▮▮

Call: ▮▮▮▮▮▮▮▮▮▮
(Call for proposals for simple programmes 2021 - Promotion of ▮▮▮▮▮▮▮)

Topic: ▮▮▮▮▮▮▮▮▮▮▮▮▮▮▮▮▮▮

Type of Action: ▮▮▮▮▮▮
▮▮▮▮▮▮▮▮▮▮

Proposal number: ▮▮▮▮▮▮▮

Proposal acronym: TEST

Type of Model Grant Agreement: ▮▮▮▮ **Action Grant Budget-Based**

Table of contents

Section	Title	Action
1	General information	Show
2	Participants	Show
3	Budget	Show

Estructura formulario parte A (Propuesta completa)

1. Información general

2. Participantes

3. Presupuesto

4. Ética y seguridad

5. Otras cuestiones

Topic:

Type of action:

Type of Model Grant Agreement:

Proposal number:

Proposal acronym:

Table of contents

Section	Title	Action
1	General information	
2	Participants	
3	Budget	
4	Ethics and security	
5	Other questions	

Example, not to complete

➢ Las partes marcadas en azul son solo de lectura

1. Información general

> Solo puede cumplimentarla los contactos del coordinador

1 – General information

Section 1 provides basic data on the proposal. It can be filled in by contacts of the coordinator. Other participants may view this section only. Read-only parts are marked in blue.

Topic	Type of action
Call	Type of Model Grant Agreement

Acronym — *Acronym is mandatory*

Proposal title — *Max 200 characters (with spaces). Must be understandable for non-specialists in your field.*

*Note that for technical reasons, the following characters are not accepted in the Proposal Title and will be removed: < > " & *

Duration in months — *Estimated duration of the project in full months.*

Fixed keyword

Fixed keyword

Free keywords — *Enter any words you think give extra detail of the scope of your proposal (max 200 characters with spaces).*

Abstract

The abstract should provide the reader with a clear understanding of the objectives of the proposal, how they will be achieved, and their relevance to the Work Programme. This summary will be used as the short description of the proposal in the evaluation process and in communications to the programme management committees and other interested parties. It must therefore be short and precise and should not contain confidential information. Use plain typed text, avoiding formulas and other special characters. If the proposal is written in a language other than English, please include an English version of this abstract in the Part B (technical description) of the proposal.

Has this proposal (or a very similar one) been submitted in the past 2 years in response to a call for proposals under any EU programme, including the current call? *A 'similar' proposal or contract is one that differs from the current one in minor ways, and in which some of the present consortium members are involved.*	○ Yes	○ No
Please give the proposal reference or contract number	XXXXX-X	

> El *abstract* se utilizará como descripción breve de la propuesta en el proceso de evaluación y en las comunicaciones con los comités de gestión del programa

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Declarations

These declarations can be filled in by any coordinator contact(s). All declarations are mandatory.

1) We declare to have the explicit consent of all applicants on their participation and on the content of this proposal.	☐
2) We confirm that the information contained in this proposal is correct and complete and that none of the project activities have started before the proposal was submitted (unless explicitly authorised in the call conditions).	☐
3) We declare: – to be fully compliant with the eligibility criteria set out in the call – not to be subject to any exclusion grounds under the EU Financial Regulation 2018/1046 – to have the financial and operational capacity to carry out the proposed project.	☐
4) We acknowledge that all communication will be made through the Funding & Tenders Portal electronic exchange system and that access and use of this system is subject to the Funding & Tenders Portal Terms & Conditions.	☐
5) We have read, understood and accepted the Funding & Tenders Portal Terms & Conditions and Privacy Statement that set out the conditions of use of the Portal and the scope, purposes, retention periods, etc. for the processing of personal data of all data subjects whose data we communicate for the purpose of the application, evaluation, award and subsequent management of our grant, prizes and contracts (including financial transactions and audits).	☐
6) We declare that the proposal complies with ethical principles (including the highest standards of research integrity as set out in the ALLEA European Code of Conduct for Research Integrity, as well as applicable international and national law, including the Charter of Fundamental Rights of the European Union and the European Convention on Human Rights and its Supplementary Protocols. Appropriate procedures, policies and structures are in place to foster responsible research practices, to prevent questionable research practices and research misconduct, and to handle allegations of breaches of the principles and standards in the Code of Conduct.	☐
7) We declare that the proposal has an exclusive focus on civil applications (activities intended to be used in military application or aiming to serve military purposes cannot be funded). If the project involves dual-use items in the sense of Regulation 2021/821, or other items for which authorisation is required, we confirm that we will comply with the applicable regulatory framework (e.g. obtain export/import licences before these items are used).	☐

8) We confirm that the activities proposed do not – aim at human cloning for reproductive purposes; – intend to modify the genetic heritage of human beings which could make such changes heritable (with the exception of research relating to cancer treatment of the gonads, which may be financed), or – intend to create human embryos solely for the purpose of research or for the purpose of stem cell procurement, including by means of somatic cell nuclear transfer. – lead to the destruction of human embryos (for example, for obtaining stem cells) These activities are excluded from funding.	☐
9) We confirm that for activities carried out outside the Union, the same activities would have been allowed in at least one EU Member State	☐
10) [Additional option for LUMP SUM Grants: For Lump Sum Grants with a detailed budget table: We understand and accept that the EU lump sum grants must be reliable proxies for the actual costs of a project and confirm that the detailed budget for the proposal has been established in accordance with our usual cost accounting practices and in compliance with the basic eligibility conditions for EU actual cost grants (see AGA — Annotated Grant Agreement, art 6) and exclude costs that are ineligible under the Programme. Purchases and subcontracting costs must be done taking into	☐

> ➢ **Es obligatorio cumplimentar las declaraciones**

2. Participantes

2 – Participants

List of participating organisations

#	Participating Organisation Legal Name	Country
1		
2		
3		

> ➢ Los datos de la organización se rellena automáticamente según la información cargada en el sistema de registro de organizaciones de la CE (*Participant Register*). No se pueden cambiar en el formulario.

Organisation data

The section shows the administrative data of the participating organisation as registered and/or validated in the central registry of organisations of the European Commission, linked to the given PIC number. Data in blue is read-only, modification is not possible in the proposal forms. For more information on how to modify this information, please visit the online manual on the participant register.

PIC	Legal name

Short name

Address of the organisation

- Street
- Town
- Postcode
- Country
- Webpage

Specific legal statuses

Read more about legal statuses

Public unknown	unknown	Legal person	
Non-profit	unknown		
International organisation	unknown		
International organisation of European interest	unknown		
Secondary or Higher education establishment	unknown		
Research organisation	unknown		

SME status

The enterprise data of the organisation is defined using the Participant Register. Changes to the self-declared or self-assessed SME data can be performed by the self-registrant or to the LEAR (Legal Entity Appointed Representative) in the Participant Register.

SME self declared status	unknown
SME self-assessment	unknown
SME validation sme	unknown

Based on the above details of the Participant Registry the organisation is not an SME (small- and medium-sized enterprise) for the call.

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2. Participantes

> ➢ Hay que indicar si hay relaciones de dependencia entre los participantes

Two participants (legal entities) are dependent on each other where there is a controlling relationship between them:

- *A legal entity is under the same direct or indirect control as another legal entity;or*

- *A legal entity directly or indirectly controls another legal entity;or*

- *A legal entity is directly or indirectly controlled by another legal entity*

> ➤ El "**main contact person**" de cada entidad, será la persona con la que los servicios de la UE contactaría en relación con la propuesta.

> ➤ Los datos (nombre, apellidos y correo electrónico) de las personas de contacto principales se introducen y modifican en el paso "Participantes" del asistente del sistema de presentación

2. Participantes

Tabla de investigadores. (Novedad HE):

➢ Con fines estadísticos para la CE para seguimiento de la carrera de los investigadores (Indicador HE)

➢ Se usará en caso de que sea necesario llegar al criterio de "*Gender balance*" en los desempates.

➢ Se incluyen los investigadores de todos los tipos de entidad

2. Participantes

Role of participating organisation in the project *Applicants may select more than one option.*	
Project management	☐
Communication, dissemination and engagement	☐
Provision of research and technology infrastructure	☐
Co-definition of research and market needs	☐
Civil society representative	☐
Policy maker or regulator, incl. standardisation body	☐
Research performer	☐
Technology developer	☐
Testing/validation of approaches and ideas	☐
Prototyping and demonstration	☐
IPR management incl. technology transfer	☐
Public procurer of results	☐
Private buyer of results	☐
Finance provider (public or private)	☐
Education and training	☐
Contributions from the social sciences or/and the humanities	☐
Other Specify (50 character limit):	☐

Tabla rol de la entidad (Novedad HE):

➤ Hay que especificar el rol de la entidad en el proyecto

➤ Una misma entidad puede tener varios roles

2. Participantes

Otras tablas con información sobre el participante (En H2020 información incluida en la sección 4 de la parte B)

List of up to 5 publications, widely-used datasets, software, goods, services, or any other achievements relevant to the call content.

Type of achievement	Short description
[Publication] [Dataset] [Software] [Good] [Service] [Other achievement]	Key elements of the achievement, including a short qualitative assessment of its impact and (where available) its digital object identifier (DOI) or other type of persistent identifier (PID). Publications, in particular journal articles, are expected to be open access. Datasets are expected to be FAIR and 'as open as possible, as closed as necessary'.

List of up to 5 most relevant previous projects or activities, connected to the subject of this proposal

Name of Project or Activity	Short description

Description of any significant infrastructure and/or any major items of technical equipment, relevant to the proposed work

Name of infrastructure or equipment	Short description

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2. Participantes

Gender equality plan

Having a gender equality plan is an eligibility criterion for Public bodies, Higher education establishments and Research organisations from Member States and Associated Countries. Be aware that if the proposal is selected, having a Gender Equality Plan will be necessary before the grant agreement signature (applicable on calls with deadlines in 2022 and beyond).

Does the organisation have a Gender Equality Plan (GEP) covering the elements listed below?	○ Yes	○ No

Minimum process-related requirements (building blocks) for a GEP

- **Publication:** formal document published on the institution's website and signed by the top management
- **Dedicated resources:** commitment of human resources and gender expertise to implement it.
- **Data collection and monitoring:** sex/gender disaggregated data on personnel (and students for establishments concerned) and annual reporting based on indicators.
- **Training:** Awareness raising/trainings on gender equality and unconscious gender biases for staff and decision-makers.

Content-wise, recommended areas to be **covered** and addressed via concrete measures and targets are:

- o work-life balance and organisational culture;
- o gender balance in leadership and decision-making;
- o gender equality in recruitment and career progression;
- o integration of the gender dimension into research and teaching content;
- o measures against gender-based violence including sexual harassment.

Plan de igualdad de género (Novedad HE):

➢ **Autodeclaración** para todos los participantes

➢ **Condición de elegibilidad** a partir de convocatorias con fechas de cierre en 2022 para entidades públicas, centros de investigación y universidades de los EEMM y PPAA.

➢ Hay que tenerlo publicado antes de la firma del GA, sino la entidad no sería eligible para recibir financiación.

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3. Presupuesto

Tabla de presupuesto (Budget table):

No	Participant name	Country	A. Personnel costs/€ (a1)	B. Subcontracting costs/€ (b)	C. Purchase costs			D. Other cost categories D.X [specific cost]	E. Indirect costs/€ (e) = 25% [(a1) + (c1) + (c2) + (c3) + (d7)]	Total eligible costs (h) = (a1) + (b) + (c1) + (c2) + (c3) +	Funding rate (U)	Maximum EU contribution to eligible costs (l) = (U) * (h)	Requested EU contribution to eligible costs/€ (Requested grant amount) (m) (n)	Income generated by the action (o)	Financial contributions (q)	Own resources (r)	Total estimated income (s)=(n)+(o)+(p)+(q)+(r)
					C.1 Travel	C.2 Equipm	C.3 Other										
1	Participant 1	NL															
2	Participant 2	LB															
	Affiliated Entity	LB															
3	Participant 3	DE															
	Associated Partner	AR															
	Total																

Estimated expenditure (Estimated eligible costs) | **Estimated income** (Requested EU contribution — EU contribution to eligible costs; Revenues; Other sources of financing)

A.- Costes de personal
B.- Costes subcontratación.
C.- Compras (viajes, equipamiento, otros bienes, trabajos y servicios)
D.- Otras categorías de costes
E.- Costes indirectos
Costes totales elegibles.

Los *Associated Partners* tienen que incluir sólo su presupuesto total en "*Other sources of funding*" (como "*financial contributions*" en caso de que recibieran financiación de su Administración (p. ej. CH) o como "*own ressources*")

4. Ética y seguridad

Tabla sobre asuntos éticos

1. HUMAN EMBRYONIC STEM CELLS AND HUMAN EMBRYOS			Page
Does this activity involve Human Embryonic Stem Cells (hESCs)?		Yes No	
If YES:	Will they be directly derived from embryos within this project?	Yes No	
	Are they previously established cells lines?	Yes No	
	Are the cell lines registered in the European registry for human embryonic stem cell lines?	Yes No	
Does this activity involve the use of human embryos?		Yes No	
If YES:	Will the activity lead to their destruction?	Yes No	
2. HUMANS			Page
Does this activity involve human participants?		Yes No	
If YES:	Are they volunteers for nonmedical studies (e.g. social or human sciences research)?	Yes No	
	Are they healthy volunteers for medical studies?	Yes No	
	Are they patients for medical studies?	Yes No	
	Are they potentally vulnerable individuals or groups?	Yes No	
	Are they children/minors?	Yes No	
	Are they other persons unable to give informed consent?	Yes No	

Example, not to complete

1. HUMAN EMBRYONIC STEM CELLS AND HUMAN EMBRYOS

2. HUMANS

3. HUMAN CELLS / TISSUES

4. PERSONAL DATA

5. ANIMALS

6. NON-EU COUNTRIES

7. ENVIRONMENT, HEALTH and SAFETY.

8. ARTIFICIAL INTELLINGENCE.

9. OTHER ETHICS ISSUES (e.g. Misuse)

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Autoevaluación ética

Si se responde "**Sí**" a alguna de las preguntas de la tabla, hay que proporcionar información adicional sobre esa cuestión de ética en el apartado de **"Ethics Self-Assesment".**

[Guía para completar la autoevaluación ética](#)

EU Grants

How to complete your ethics self-assessment

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Cuestionario sobre seguridad   Novedad HE

Escrutinio de seguridad sistemática en todas las propuestas basada en la autoevaluación incluida en la propuesta y centrada en determinar:

- ➤ Si la propuesta utiliza o genera información clasificada de la UE.

- ➤ El potencial de uso indebido de los resultados (que podrían canalizarse hacia la delincuencia o el terrorismo).

- ➤ Si las actividades implican información o materiales sujetos a restricciones de seguridad nacional.

Security issues table

Please indicate, by answering Yes or No to all of the questions in the below table, if the proposed activity will use and/or generate information which might raise security concerns. If an answer is Yes, then indicate in the adjacent box at which page in your full proposal further information relating to that issue can be found.

1. EU classified information (EUCI)[2]			Page
Does this activity involve information and/or materials requiring protection against unauthorised disclosure (EUCI)?		○ Yes ○ No	
If YES:	Is the activity going to use classified information as background[3] information?	○ Yes ○ No	
	Is the activity going to generate EU classified foreground[4] information as results?	○ Yes ○ No	
Does this activity involve non-EU countries?		○ Yes ○ No	
If YES:	Do participants from non-EU countries need to have access to EUCI?	○ Yes ○ No	
	Do the non-EU countries concerned have a security of information agreement with the EU	○ Yes ○ No	
2. MISUSE			Page
Does this activity have the potential for misuse of results?		○ Yes ○ No	
If YES:	Does the activity provide knowledge, materials and technologies that could be channelled into crime and/or terrorism?	○ Yes ○ No	
	Could the activity result in the development of chemical, biological, radiological or nuclear (CBRN) weapons and the means for their delivery?	○ Yes ○ No	
3. OTHER SECURITY ISSUES			Page
Does this activity involve information and/or materials subject to national security restrictions?		○ Yes ○ No	
If yes, please specify: *(Maximum number of characters allowed: 1000)*			

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5. Otras cuestiones

5 – Other questions

Two-stage calls

The full stage-2 proposal must be consistent with the short outline proposal submitted to the stage 1 – in particular with respect to the proposal characteristics addressing the concepts of excellence and impact.

Are there substantial differences compared to the stage-1 proposal?	○ Yes	○ No

Questions showed only in answer is Yes:
Please list the substantial differences, and indicate the reasons

☐	Partnership	List the substantial differences and indicate the reasons
☐	Budget	List the substantial differences and indicate the reasons
☐	Approach	List the substantial differences and indicate the reasons

> Para convocatorias en dos fases, habría que indicar si ha habido cambios substanciales de la fase 1 a la fase 2.

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5. Otras cuestiones

[Calls with clinical trials: Essential information to be provided for proposals including clinical trials / studies / investigations

Clinical study means, for the purpose of this document, any systematic prospective or retrospective collection and analysis of health data obtained from individual patients or healthy persons in order to address scientific questions related to the understanding, prevention, diagnosis, monitoring or treatment of a disease, mental illness, or physical condition. It includes but it is not limited to clinical studies as defined by Regulation 536/2014 (on medicinal products), clinical investigation and clinical evaluation as defined by Regulation 2017/745 (on medical devices), performance study and performance evaluation as defined by Regulation 2017/746 (on in vitro diagnostic medical devices.

Are clinical studies / trials / investigations included in the work plan of this project?	○ Yes ○ No

Please upload the dedicated annex 'Essential information for clinical studies / trials / investigations' (a Word template is provided under 'download templates' in the up-load section for Part B and Annexes).

This document should include the relevant information of each clinical study / trial / investigation included in the work plan of this project.

Please give a short title, an acronym or a unique identifier to each clinical study / trial / investigation, to be used as a reference / identifier in the other parts of the proposal	Add Remove

➢ Cumplimentar en caso de que la propuesta implique estudios clínicos.

➢ Los estudios clínicos se subirán en la parte de Anexos

Plantilla propuestas "stage 1" de convocatorias "two-stage"

Formulario PARTE A

1- Información general

2- Participantes: *organisation data, departments carrying out de proposed work, links with other participants , main contact person*

3- Presupuesto **= TOTAL requested EU contribution (EUR)**

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