



Horizon Europe Programme

Standard Application Form  
(HE EIC Support for Ukrainian tech SMEs and  
startups)

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

Version 1.0  
03 July 2025



Application form (Part A)

Example, not to be completed



# Horizon Europe Programme

## Standard Application Form (HE EIC Support for Ukrainian tech SMEs and startups)

Application form (Part A)

Version 1.2  
12 April 2024

### Disclaimer

This document is aimed at informing potential applicants for Horizon Europe funding. It serves only as an example. The actual Web forms and templates, provided in the online submission system under the Funding and Tenders Portal, might differ from this example.

## Structure of the Proposal

The proposal contains two parts:

- **Part A** of the proposal is generated by the IT system. It is based on the information entered by the participants through the submission system in the Funding & Tenders Portal. The participants can update the information in the submission system at any time before final submission.
- **Part B** of the proposal is the narrative part that includes three sections that each correspond to an evaluation criterion. Part B needs to be uploaded as a PDF document following the templates downloaded by the applicants in the submission system for the specific call or topic. The templates for a specific call may slightly differ from the example provided in this document.

The electronic submission system is an online wizard that guides you step-by-step through the preparation of your proposal. The submission process consists of 6 steps:

- Step 1: Logging in the Portal
- Step 2: Select the call, topic and type of action in the Portal
- Step 3: Create a draft proposal: Title, acronym, summary, main organisation and contact details
- Step 4: Manage your parties and contact details: add your partner organisations and contact details.
- Step 5: Edit and complete web forms for proposal part A and upload proposal part B
- Step 6: Submit the proposal

- Instructions and footnotes in green will not appear in the text generated by the IT system.
- For options [in square brackets]: the option that applies will be automatically shown in the IT system (Part A) or included in the template of Part B offered by the IT system or you must select the appropriate value from a predefined list.
- For fields in [grey in square brackets] (even if they are part of an option as specified in the previous item): enter the appropriate data in the IT system.
- Data in coloured fields will be prefilled by the IT tool.

HISTORY OF CHANGES		
Version	Publication date	Changes
1.0	15.04.2021	▪ Initial version.
1.1	10.02.2022	▪ Added definitions for role of participants
1.2	12.04.2024	▪ Adapted budget table to lump sum funding

### Disclaimer

This document is aimed at informing potential applicants for Horizon Europe funding. It serves only as an example. The actual Web forms and templates, provided in the online submission system under the Funding and Tenders Portal, might differ from this example.

# Horizon Europe

## Application forms (Part A)

**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	

*The forms must be filled in for each proposal using the templates available in the Submission System. Some data fields in the forms are pre-filled based on the previous steps in the Submission wizard.*

Proposal ID XXXXXXXXXX	Acronym XXXXXXXX	Participant short name: XXXX

## 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	<i>Acronym is mandatory</i>
Proposal title	<i>Max 200 characters (with spaces). Must be understandable for non-specialists in your field.</i>
	<i>Note that for technical reasons, the following characters are not accepted in the Proposal Title and will be removed: &lt; &gt; " &amp;</i>
Duration in months	<i>Estimated duration of the project in full months.</i>
Fixed keywords	
	<i>Note that for this call, applicants have to select minimum 3 and maximum 6 fixed keywords.</i>
Free keywords	<i>Enter any words you think give extra detail of the scope of your proposal (max 200 characters with spaces).</i>

### 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? <i>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.</i>	<input type="radio"/> Yes	<input type="radio"/> No
Please give the proposal reference or contract number	XXXXX-X	

## 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.	<input type="checkbox"/>
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).	<input type="checkbox"/>
3) We declare: <ul style="list-style-type: none"> <li>– to be fully compliant with the eligibility criteria set out in the call</li> <li>– not to be subject to any exclusion grounds under the <a href="#">EU Financial Regulation 2018/1046</a></li> <li>– to have the financial and operational capacity to carry out the proposed project.</li> </ul>	<input type="checkbox"/>
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 <a href="#">Funding &amp; Tenders Portal Terms &amp; Conditions</a> .	<input type="checkbox"/>
5) We have read, understood and accepted the <a href="#">Funding &amp; Tenders Portal Terms &amp; Conditions</a> and <a href="#">Privacy Statement</a> 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).	<input type="checkbox"/>
6) We declare that the proposal complies with ethical principles (including the highest standards of research integrity as set out in the <a href="#">ALLEA European Code of Conduct for Research Integrity</a> , 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. <a href="#">Appropriate procedures, policies and structures</a> 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.	<input type="checkbox"/>
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 <a href="#">Regulation 2021/821</a> , 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).	<input type="checkbox"/>
8) We confirm that the activities proposed do not <ul style="list-style-type: none"> <li>– aim at human cloning for reproductive purposes;</li> <li>– 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</li> <li>– 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.</li> <li>– lead to the destruction of human embryos (for example, for obtaining stem cells)</li> </ul> <p>These activities are excluded from funding.</p>	<input type="checkbox"/>
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.	<input type="checkbox"/>
10) <i>[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 <a href="#">AGA — Annotated Grant Agreement, art 6</a>) and exclude costs that are ineligible under the Programme. Purchases and subcontracting costs must be done taking into account best value for money and must be free of conflict of interest. ]</i>	<input type="checkbox"/>

Proposal ID XXXXXXXXXX	Acronym XXXXXXXX	Participant short name: XXXX

The coordinator is only responsible for the information relating to their own organisation. Each applicant remains responsible for the information declared for their organisation. If the proposal is retained for EU funding, they will all be required to sign a declaration of honour.

**False statements** or incorrect information may lead to administrative sanctions under the EU Financial Regulation.

Example, not to be completed

## 2 – Participants

### List of participating organisations

#	Participating Organisation Legal Name	Country
1		
2		
3		

Coordinator contacts have the rights to:

- add, delete, edit and re-order partners in the consortium
- add, delete, edit and re-order contact points for those organisations
- edit all sections of the administrative forms
- upload, delete, view and download Part B and Annexes (when required for the call)
- submit the proposal

Participant contacts may:

- view all the information in this screen, but not edit it
- edit only the section for their organisation in the administrative forms (including budget)
- view the entire administrative forms
- view/download the Part B and other Annexes

You can manage the list of organisations and access rights of persons at Step 4 of the submission process. You may identify and give access to as many contact persons of the selected organisations as you wish. The identification is based upon the e-mail address of the person. When you add a contact person, you will be prompted to supply the contact details: name, e-mail, phone.

Person in charge of the proposal (main contact person): Each organisation needs to have one main contact person identified; the main contact person will have to fill in full contact details in the administrative form. The 'Main Contact Person' for the coordinating organisation (Participant no. 1) will become the primary contact person for the Services. Other contact persons may also be identified and may receive read-only or full access rights. Contact persons with full access rights of the coordinator (Participant no. 1) will be called 'Coordinator contacts' in the Funding & Tenders Portal, while for the other participants 'Participant Contacts'; contact persons with read-only rights will be called 'Team Members'. Other contact persons are listed with basic details in the administrative form.

Access rights: The main contact person and contact persons of the coordinator with full access rights have the same level of rights: they can manage the list of participants and contacts, edit any part of the administrative part of the proposal and upload any attachments (eg. Part B - technical description), and submit the proposal. Contact persons with read-only rights can only view/download the information. Participant contacts with full access rights can only edit their section of the administrative form and view all proposal data. Access rights can be revoked by the Coordinating Organisation contacts. The person who created the proposal cannot be deleted.

Invitation: All contacts will receive an e-mail and a notification to the Portal about the invitation to the proposal upon saving the data at Step 4 of the submission process.

## 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
<i>Short name</i>	
<i>Address of the organisation</i>	
Street	
Town	
Postcode	
Country	
Webpage	
<i>Specific legal statuses</i>	
<a href="#">Read more about legal statuses.</a>	
Public ..... unknown	Legal person .....
unknown	
Non-profit..... unknown	
International organisation ..... unknown	
International organisation of European interest ..... unknown	
Secondary or Higher education establishment ..... unknown	
Research organisation ..... unknown	
<i>SME status</i>	
<i>The enterprise data of the organisation is taken from the Participant Register. Changes to the self-declared or self-assessed SME data can be performed by the self-registrant or by the LEAR (Legal Entity Appointed Representative) in the Participant Register.</i>	
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.	

### Departments carrying out the proposed work

The information serves mainly statistical purposes. For determining the eligibility of the proposal, the official address of the organisation is taken into account.

#### Department 1

Department name	<input type="text"/>	<input type="checkbox"/> not applicable
	<input type="checkbox"/> Same as organisation address	
Street	<input type="text" value="Please enter street name and number"/>	
Town	<input type="text"/>	<input type="text"/>
Postcode	<input type="text"/>	<input type="text"/>
Country	<input type="text"/>	<input type="text"/>

### Links with other participants

Please indicate if there are dependencies with other participants of the proposal.

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. Control:

Legal entity A controls legal entity B if:

\* A, directly or indirectly, holds more than 50% of the nominal value of the issued share capital or a majority of the voting rights of the shareholders or associates of B, or

\* A, directly or indirectly, holds in fact or in law the decision-making powers in B.

The following relationships between legal entities shall not in themselves be deemed to constitute controlling relationships:

(a) the same public investment corporation, institutional investor or venture-capital company has a direct or indirect holding of more than 50 % of the nominal value of the issued share capital or a majority of voting rights of the shareholders or associates;

(b) the legal entities concerned are owned or supervised by the same public body.

Type of link	Participant
[Same group] [Controls] [Is controlled by]	Select one participant from the list of participants

### Main contact person

It is the main scientist or team leader in charge of the proposal for the participant. For participant number 1 (the coordinator), this will be the person the EU services will contact concerning this proposal (e.g. for additional information, invitation to hearings, sending of evaluation results, convocation to start grant agreement preparation). The data in blue is read-only. Details (name, first name and e-mail) of Main Contact persons should be edited in Step 4 of the Submission wizard.

Title Gender  Woman  Man  Non binary

First name

Last name

E-mail

Position in org.

*Please indicate the position of the person*

Department

Same as organisation

Same as organisation address

Street

Town

Post code

Country

Website

Phone 1

Phone 2

### Other contact persons

First name	Last name	e-mail	Phone

Application Forms

Proposal ID XXXXXXXXXX

Acronym XXXXXXXX

Participant short name: XXXX

**Researchers involved in the proposal**

*Include only the researchers involved in the proposal, (see below definition of ‘researcher’). You do not need to include in the table the identity of other persons involved in the proposal who are not researchers.*

*‘Researchers are professionals engaged in the conception or creation of new knowledge. They conduct research and improve or develop concepts, theories, models, techniques instrumentation, software or operational methods. (Frascati Manual 2015)’*

*Include also person in charge of the proposal if a researcher*

Title	First Name	Last Name	Gender	Nationality	E-mail	Career stage <sup>1</sup>	Role of researcher (in the project)	Reference Identifier	Type of identifier
			[Woman] [Man] [Non-binary]			[Category A – Top grade researcher] [Category B – Senior researcher] [Category C – Recognised researcher] [Category D – First stage researcher]	[Leading] [Team member]		[ORCID] [Researcher Id] [Other - specify]

<sup>1</sup> Career stages as defined in Frascati 2015 manual:

Category A – Top grade researcher: the single highest grade/post at which research is normally conducted. Example: ‘Full professor’ or ‘Director of research’.

Category B – Senior researcher: Researchers working in positions not as senior as top position but more senior than newly qualified doctoral graduates (ISCED level 8). Examples: ‘associate professor’ or ‘senior researcher’ or ‘principal investigator’.

Category C – Recognised researcher: the first grade/post into which a newly qualified doctoral graduate would normally be recruited. Examples: ‘assistant professor’, ‘investigator’ or ‘post-doctoral fellow’.

Category D – First stage researcher: Either doctoral students at the ISCED level 8 who are engaged as researchers, or researchers working in posts that do not normally require a doctorate degree. Examples: ‘PhD students’ or ‘junior researchers’ (without a PhD).

<b>Role of participating organisation in the project</b> <i>Applicants may select more than one option.</i>		<b>Definitions</b>
Project management	<input type="checkbox"/>	Click if your organisation will do project management activities (i.e. assigning the tasks, reporting and interface with the EC). These tasks are normally carried out by the coordinator, but other participants can also contribute.
Communication, dissemination and engagement	<input type="checkbox"/>	Click if your organisation will be in charge of communication, dissemination and engagement. This can be centralised by one partner or split across the partners.
Provision of research and technology infrastructure	<input type="checkbox"/>	Click if your organisation is providing a research facility or research equipment.
Co-definition of research and market needs	<input type="checkbox"/>	Click if your organisation will be involved in the co-defining the research and market needs. Usually it is a company that intends to later use the research results, or a NGO that will use the solution. This will help the project further tailor its results to respond to specific needs of the end user.
Civil society representative	<input type="checkbox"/>	Click if your organisation belongs to civil society (NGO, association, organisation, consumer group, community group, charity, etc.).
Policy maker or regulator, incl. standardisation body	<input type="checkbox"/>	Click if your organisation is a policy maker (local, regional, national, European level), regulator or a standardisation body.
Research performer	<input type="checkbox"/>	Click if your organisation is in charge of performing the research during the project.
Technology developer	<input type="checkbox"/>	Click if your organisation is in charge of developing the technology during or after the project.
Testing/validation of approaches and ideas	<input type="checkbox"/>	Click if your organisation is in charge of testing/validating the approach and ideas.
Prototyping and demonstration	<input type="checkbox"/>	Click if your organisation is in charge of developing the prototypes and performing demonstrations.
IPR management incl. technology transfer	<input type="checkbox"/>	Click if your organisation is in charge of IPR management including technology transfer at the end of the grant.
Public procurer of results	<input type="checkbox"/>	Click if your organisation (public authority, hospital, university, local government, etc) will be using the results afterwards.
Private buyer of results	<input type="checkbox"/>	Click if your organisation (from the private sector) will be using the results afterwards.
Finance provider (public or private)	<input type="checkbox"/>	Click if your organisation will be providing the financing for the exploitation during or after the end of the project.
Education and training	<input type="checkbox"/>	Click if your organisation is in charge of educating and training researchers.
Contributions from the social sciences or/and the humanities	<input type="checkbox"/>	Click if your organisation is in charge of contributing to the social sciences or/and the humanities dimension to the research project.
Other Specify (50 character limit):	<input type="checkbox"/>	

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

<b>Type of achievement</b>	<b>Short description</b>
[Publication] [Dataset] [Software]	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'.

Application Forms

Proposal ID XXXXXXXXXX

Acronym XXXXXXXX

Participant short name: XXXX

[Good]	
[Service]	
[Other achievement]	

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

## 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:

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

Example, not to be completed

### 3 – Budget for the proposal

---

No	Name of Beneficiary	Country	Role	Requested grant amount
1				
Total				

Example, not to be completed

## 4 – Ethics and Security

### Ethics issues table

This table should be completed as an essential part of your proposal. Please go through the table and indicate which elements concern your proposal by answering 'Yes' or 'No'. If you answer 'Yes' to any of the questions,

- indicate in the adjacent box at which page in your technical description further information relating to that ethics issue can be found, and
- provide additional information on that ethics issue in the Ethics Self-Assessment section.

For more information on each of the ethics issues and how to address them, including detailed legal references, see the guidelines '[How to Complete your Ethics Self-Assessment](#)'.

1. HUMAN EMBRYONIC STEM CELLS AND HUMAN EMBRYOS			Page
Does this activity involve Human Embryonic Stem Cells (hESCs)?		<input type="radio"/> Yes <input type="radio"/> No	
If YES:	Will they be directly derived from embryos within this project?	<input type="radio"/> Yes <input type="radio"/> No	
	Are they previously established cells lines?	<input type="radio"/> Yes <input type="radio"/> No	
	Are the cell lines registered in the European registry for human embryonic stem cell lines?	<input type="radio"/> Yes <input type="radio"/> No	
Does this activity involve the use of human embryos?		<input type="radio"/> Yes <input type="radio"/> No	
If YES:	Will the activity lead to their destruction?	<input type="radio"/> Yes <input type="radio"/> No	
2. HUMANS			Page
Does this activity involve human participants?		<input type="radio"/> Yes <input type="radio"/> No	
If YES:	Are they volunteers for non medical studies (e.g. social or human sciences research)?	<input type="radio"/> Yes <input type="radio"/> No	
	Are they healthy volunteers for medical studies?	<input type="radio"/> Yes <input type="radio"/> No	
	Are they patients for medical studies?	<input type="radio"/> Yes <input type="radio"/> No	
	Are they potentially vulnerable individuals or groups?	<input type="radio"/> Yes <input type="radio"/> No	
	Are they children/minors?	<input type="radio"/> Yes <input type="radio"/> No	
	Are they other persons unable to give informed consent?	<input type="radio"/> Yes <input type="radio"/> No	
Does this activity involve interventions (physical also including imaging technology, behavioural treatments, etc.) on the study participants?		<input type="radio"/> Yes <input type="radio"/> No	
If YES:	Does it involve invasive techniques?	<input type="radio"/> Yes <input type="radio"/> No	
	Does it involve collection of biological samples?	<input type="radio"/> Yes <input type="radio"/> No	

Application Forms		
Proposal ID XXXXXXXXXX	Acronym XXXXXXXX	Participant short name: XXXX
Does this activity involve conducting a clinical study as defined by the Clinical Trial Regulation (EU 536/2014)? (using pharmaceuticals, biologicals, radiopharmaceuticals, or advanced therapy medicinal products)		<input type="radio"/> Yes <input type="radio"/> No
If YES:	Is it a clinical trial?	<input type="radio"/> Yes <input type="radio"/> No
	Is it a low-intervention clinical trial?	<input type="radio"/> Yes <input type="radio"/> No
3. HUMAN CELLS / TISSUES (not covered by section 1)		Page
Does this activity involve the use of human cells or tissues?		<input type="radio"/> Yes <input type="radio"/> No
If YES:	Are they human embryonic or foetal cells or tissues?	<input type="radio"/> Yes <input type="radio"/> No
	Are they available commercially?	<input type="radio"/> Yes <input type="radio"/> No
	Are they obtained within this project?	<input type="radio"/> Yes <input type="radio"/> No
	Are they obtained from another project, laboratory or institution?	<input type="radio"/> Yes <input type="radio"/> No
	Are they obtained from biobank?	<input type="radio"/> Yes <input type="radio"/> No
4. PERSONAL DATA		Page
Does this activity involve processing of personal data?		<input type="radio"/> Yes <input type="radio"/> No
If YES:	Does it involve the processing of special categories of personal data (e.g.: sexual lifestyle, ethnicity, genetic, biometric and health data, political opinion, religious or philosophical beliefs)?	<input type="radio"/> Yes <input type="radio"/> No
	If YES: Does it involve processing of genetic, biometric or health data?	<input type="radio"/> Yes <input type="radio"/> No
	Does it involve profiling, systematic monitoring of individuals, or processing of large scale of special categories of data or intrusive methods of data processing (such as, surveillance, geolocation tracking etc.)?	<input type="radio"/> Yes <input type="radio"/> No
Does this activity involve further processing of previously collected personal data (including use of preexisting data sets or sources, merging existing data sets)?		<input type="radio"/> Yes <input type="radio"/> No
Is it planned to export personal data from the EU to non-EU countries?		<input type="radio"/> Yes <input type="radio"/> No
If YES:	Specify the type of personal data and countries involved:	
Is it planned to import personal data from non-EU countries into the EU or from a non-EU country to another non-EU country?		<input type="radio"/> Yes <input type="radio"/> No
If YES:	Specify the type of personal data and countries involved	

Application Forms		
Proposal ID XXXXXXXXXX	Acronym XXXXXXXX	Participant short name: XXXX
Does this activity involve the processing of personal data related to criminal convictions or offences?		<input type="radio"/> Yes <input type="radio"/> No
<b>5. ANIMALS</b>		<b>Page</b>
Does this activity involve animals?		<input type="radio"/> Yes <input type="radio"/> No
If <b>YES</b> :	Are they vertebrates?	<input type="radio"/> Yes <input type="radio"/> No
	Are they non-human primates (NHP)?	<input type="radio"/> Yes <input type="radio"/> No
	Are they genetically modified?	<input type="radio"/> Yes <input type="radio"/> No
	Are they cloned farm animals?	<input type="radio"/> Yes <input type="radio"/> No
	Are they endangered species?	<input type="radio"/> Yes <input type="radio"/> No
<b>6. NON-EU COUNTRIES</b>		<b>Page</b>
Will some of the activities be carried out in non-EU countries?		<input type="radio"/> Yes <input type="radio"/> No
If <b>YES</b> :	Specify the countries:	
In case non-EU countries are involved, do the activities undertaken in these countries raise potential ethics issues?		<input type="radio"/> Yes <input type="radio"/> No
If <b>YES</b> :	Specify the countries:	
Is it planned to use local resources (e.g. animal and/or human tissue samples, genetic material, live animals, human remains, materials of historical value, endangered fauna or flora samples, etc.)?		<input type="radio"/> Yes <input type="radio"/> No
Is it planned to import any material from non-EU countries into the EU or from a non-EU country to another non-EU country? For data imports, see section 4.		<input type="radio"/> Yes <input type="radio"/> No
If <b>YES</b> :	Specify material and countries involved:	
Is it planned to export any material from the EU to non-EU countries?		<input type="radio"/> Yes <input type="radio"/> No
If <b>YES</b> :	Specify material and countries involved:	
Does this activity involves <a href="#">low and/or lower-middle income countries</a> ? (if yes, detail the benefit-sharing actions planned in the self-assessment)		<input type="radio"/> Yes <input type="radio"/> No
Could the situation in the country put the individuals taking part in the activity at risk?		<input type="radio"/> Yes <input type="radio"/> No
<b>7. ENVIRONMENT, HEALTH and SAFETY</b>		<b>Page</b>

Application Forms		
Proposal ID XXXXXXXXXX	Acronym XXXXXXXX	Participant short name: XXXX
Does this activity involve the use of substances or processes that may cause harm to the environment, to animals or plants (during the implementation of the activity or further to the use of the results, as a possible impact)?	<input type="radio"/> Yes <input type="radio"/> No	
Does this activity deal with endangered fauna and/or flora / protected areas?	<input type="radio"/> Yes <input type="radio"/> No	
Does this activity involve the use of substances or processes that may cause harm to humans, including those performing them (during the implementation of the activity or further to the use of the results, as a possible impact)?	<input type="radio"/> Yes <input type="radio"/> No	
<b>8. ARTIFICIAL INTELLIGENCE</b>		Page
Does this activity involve the development, deployment and/or use of Artificial Intelligence based systems? (if yes, detail in the self-assessment whether that could raise ethical concerns related to human rights and values and detail how this will be addressed).	<input type="radio"/> Yes <input type="radio"/> No	
<b>9. OTHER ETHICS ISSUES</b>		Page
Are there any other ethics issues that should be taken into consideration?	<input type="radio"/> Yes <input type="radio"/> No	
<i>Please specify: (Maximum number of characters allowed: 1000)</i>		

I confirm that I have taken into account all ethics issues above and that, if any ethics issues apply, I will complete the ethics self-assessment as described in the guidelines [‘How to Complete your Ethics Self-Assessment’](#).

Example, not to be completed

## ETHICS SELF-ASSESSMENT

If you have entered any issues in the ethics issue table, you must perform an ethics self-assessment in accordance with the guidelines "[How to Complete your Ethics Self-Assessment](#)" and complete the table below.

### Ethical dimension of the objectives, methodology and likely impact

Explain in detail the identified issues in relation to:

- objectives of the activities (e.g. study of vulnerable populations, etc.)
- methodology (e.g. clinical trials, involvement of children, protection of personal data, etc.)
- the potential impact of the activities (e.g. environmental damage, stigmatisation of particular social groups, political or financial adverse consequences, misuse, etc.)

### Compliance with ethical principles and relevant legislations

Describe how the issue(s) identified in the ethics issues table above will be addressed in order to adhere to the ethical principles and what will be done to ensure that the activities are compliant with the E U/ national legal and ethical requirements of the country or countries where the tasks are to be carried out. It is reminded that for **activities performed in a non-EU countries**, they should also be allowed in at least one EU Member State.

## Application Forms

Proposal ID **XXXXXXXXXX**

Acronym **XXXXXXXX**

Participant short name: **XXXX**

### Security issues table

Please go through the table and indicate which elements concern your proposal by answering YES or NO.

If you answer YES to any of the questions:

- indicate in the adjacent box at which page in your full proposal further information relating to that security issue can be found, and
- provide additional information on this security issue in the Security self-assessment section below.

For more information on potential security issues and how to address them, see the guidance [How to handle security-sensitive projects](#) and the programme-specific guidelines [Classification of information in Horizon Europe projects](#).

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

<sup>2</sup> According to the Commission Decision (EU, Euratom) 2015/444 of 13 March 2015 on the security rules for protecting EU classified information, “European Union classified information (EUCI) means any information or material designated by an EU security classification, the unauthorised disclosure of which could cause varying degrees of prejudice to the interests of the European Union or of one or more of the Member States”.

<sup>3</sup> Classified background information is information that is already classified by a country and/or international organisation and/or the EU and is going to be used by the project. In this case, the project must have in advance the authorisation from the originator of the classified information, which is the entity (EU institution, EU Member State, third state or international organisation) under whose authority the classified information has been generated.

<sup>4</sup> EU classified foreground information is information (documents/deliverables/materials) planned to be generated by the project and that needs to be protected from unauthorised disclosure. The originator of the EUCI generated by the project is the European Commission.

Application Forms		
Proposal ID XXXXXXXXXX	Acronym XXXXXXXX	Participant short name: XXXX
If yes, please specify: (Maximum number of characters allowed: 1000)		
SECURITY SELF-ASSESSMENT		
If you have answered YES for one or more of the questions indicated above, describe the measures you intend to take to solve/avoid them. For more information, see the guidelines <a href="#">Classification of information in Horizon Europe projects</a> , <a href="#">Classification of information in Digital Europe projects</a> , <a href="#">Classification of information in EDF projects</a> .		
Please specify (Maximum number of characters allowed: 5000)		

## 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?	<input type="radio"/> Yes	<input type="radio"/> No
---	---------------------------	--------------------------

Questions showed only in answer is Yes:

Please list the substantial differences, and indicate the reasons

<input type="checkbox"/>	Partnership	List the substantial differences and indicate the reasons
<input type="checkbox"/>	Budget	List the substantial differences and indicate the reasons
<input type="checkbox"/>	Approach	List the substantial differences and indicate the reasons

### [Additional modular extension for 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?	<input type="radio"/> Yes	<input type="radio"/> 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).

Version of template used	Page 20 of 21	Last saved dd/mm/yyyy HH:mm
--------------------------	---------------	-----------------------------

This proposal version was submitted by [Name, FAMILY NAME] on [dd/mm/yyyy HH:mm:ss] Brussels Local Time. Issued by the Funding and Tenders Portal Submission Service

Application Forms

Proposal ID XXXXXXXXXX

Acronym XXXXXXXX

Participant short name: XXXX

*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

]

Example, not to be completed

## Project proposal – Technical description (Part B)

Example, not to be completed



# Horizon Europe Programme

## Standard Application Form (HE EIC Support for Ukrainian tech SMEs and startups)

Project proposal – Technical description (Part B)

Version 1.0  
03 July 2025



**Structure of the Proposal**

The proposal contains two parts:

- **Part A** of the proposal is generated by the IT system. It is based on the information entered by the participants through the submission system in the Funding & Tenders Portal. The participants can update the information in the submission system at any time before final submission.
- **Part B** of the proposal is the narrative part that includes three sections that each correspond to an evaluation criterion. Part B needs to be uploaded as a PDF document following the templates downloaded by the applicants in the submission system for the specific call or topic. The templates for a specific call may slightly differ from the example provided in this document.

The electronic submission system is an online wizard that guides you step-by-step through the preparation of your proposal. The submission process consists of 6 steps:

- Step 1: Logging in the Portal
- Step 2: Select the call, topic and type of action in the Portal
- Step 3: Create a draft proposal: Title, acronym, summary, main organisation and contact details
- Step 4: Manage your parties and contact details: add your partner organisations and contact details.
- Step 5: Edit and complete web forms for proposal part A and upload proposal part B
- Step 6: Submit the proposal

HISTORY OF CHANGES		
Version	Publication date	Changes
1.0	03.07.2025	▪ Initial version.

Example, not to be completed

## Proposal template Part B: technical description

The structure of this template must be followed when preparing your proposal. It has been designed to ensure that the important aspects of your planned work are presented in a way that will enable the experts to make an effective assessment against the evaluation criteria. Sections 1, 2 and 3 each correspond to an evaluation criterion.

Please be aware that proposals will be evaluated as they were submitted, rather than on their potential if certain changes were to be made. This means that only proposals that successfully address all the required aspects will have a chance of being funded. There will be no possibility for significant changes to content, budget and consortium composition during grant preparation.

**⚠ Page limit:** The title, list of participants and sections 1, 2 and 3, together, should not be longer than 22 pages. All tables, figures, references and any other element pertaining to these sections must be included as an integral part of these sections and are thus counted against this page limit. The number of pages included in each section of this template is only **indicative**.

The cover page (including the Seal of Excellence section, if applicable) must be completed and maintained. If missing, the proposal could be considered incomplete and therefore inadmissible.

The page limit will be applied automatically; therefore you must remove this instruction page before submitting. Remove also the table with the definition of terms and the help text added after each section.

If you attempt to upload a proposal longer than the specified limit before the deadline, you will receive an automatic warning and will be advised to shorten and re-upload the proposal. After the deadline, excess pages (in over-long proposals/applications) will be automatically made invisible, and will not be taken into consideration by the experts. The proposal is a self-contained document, meaning that experts will be instructed to ignore hyperlinks to information that is specifically designed to expand the proposal, thus circumventing the page limit.

Please, do not consider the page limit as a target! It is in your interest to keep your text as concise as possible, since experts rarely view unnecessarily long proposals in a positive light.

**⚠** The following formatting conditions apply.

The reference font for the body text of proposals is Times New Roman (Windows platforms), Times/Times New Roman (Apple platforms) or Nimbus Roman No. 9 L (Linux distributions).

The use of a different font for the body text is not advised and is subject to the cumulative conditions that the font is legible and that its use does not significantly shorten the representation of the proposal in number of pages compared to using the reference font (for example with a view to bypass the page limit).

The minimum font size allowed is 11 points. Standard character spacing and a minimum of single line spacing is to be used. This applies to the body text, including text in tables.

Text elements other than the body text, such as headers, foot/end notes, captions, formula's, may deviate, but must be legible.

The page size is A4, and all margins (top, bottom, left, right) should be at least 15 mm (not including any footers or headers).

This document is tagged. Do not delete the tags; they are needed for our internal processing of information, mostly for statistical gathering. In that light, please do not move, delete, re-order, alter tags in any way, as they might create problems in our internal processing tools. Tags do not affect or influence the outcome of your application.

<b>DEFINITIONS</b>	
<b>Critical risk</b>	<p>A critical risk is a plausible event or issue that could have a high adverse impact on the ability of the project to achieve its objectives.</p> <p>Level of likelihood to occur (Low/medium/high): The likelihood is the estimated probability that the risk will materialise even after taking account of the mitigating measures put in place.</p> <p>Level of severity (Low/medium/high): The relative seriousness of the risk and the significance of its effect.</p>
<b>Deliverable</b>	<p>A report that is sent to the Commission or Agency providing information to ensure effective monitoring of the project. There are different types of deliverables (e.g. a report on specific activities or results, data management plans, ethics or security requirements).</p>
<b>Impacts</b>	<p>Wider long term effects on society (including the environment), the economy and science, enabled by the outcomes of R&amp;I investments (long term). Impacts generally occur some time after the end of the project.</p> <p><i>Example: The deployment of the advanced forecasting system enables each airport to increase maximum passenger capacity by 15% and passenger average throughput by 10%, leading to a 28% reduction in infrastructure expansion costs.</i></p>
<b>Milestone</b>	<p>Control points in the project that help to chart progress. Milestones may correspond to the achievement of a key result, allowing the next phase of the work to begin. They may also be needed at intermediary points so that, if problems have arisen, corrective measures can be taken. A milestone may be a critical decision point in the project where, for example, the consortium must decide which of several technologies to adopt for further development. The achievement of a milestone should be verifiable.</p>
<b>Objectives</b>	<p>The goals of the work performed within the project, in terms of its research and innovation content. This will be translated into the project's results. These may range from tackling specific research questions, demonstrating the feasibility of an innovation, sharing knowledge among stakeholders on specific issues. The nature of the objectives will depend on the type of action, and the scope of the topic.</p>
<b>Outcomes</b>	<p>The expected effects, over the medium term, of projects supported under a given topic. The results of a project should contribute to these outcomes, fostered in particular by the dissemination and exploitation measures. This may include the uptake, diffusion, deployment, and/or use of the project's results by direct target groups. Outcomes generally occur during or shortly after the end of the project.</p> <p><i>Example: 9 European airports adopt the advanced forecasting system demonstrated during the project.</i></p>
<b>Research output</b>	<p>Results generated by the action to which access can be given in the form of scientific publications, data or other engineered outcomes and processes such as software, algorithms, protocols and electronic notebooks.</p>
<b>Results</b>	<p>What is generated during the project implementation. This may include, for example, know-how, innovative solutions, algorithms, proof of feasibility, new business models, new business plans, prototypes, demonstrators, databases and datasets, , networks, etc. Most project results (inventions, scientific works, etc.) are 'Intellectual Property', which may, if appropriate, be protected by formal 'Intellectual Property Rights'.</p> <p><i>Example: Successful large-scale demonstrator: trial with 3 airports of an advanced forecasting</i></p>

	<i>system for proactive airport passenger flow management.</i>
<b>Technology Readiness Level</b>	See EIC Work Programme under Glossary section. <a href="https://euraxess.ec.europa.eu/career-development/researchers/manual-scientific-entrepreneurship/major-steps/trl">TRL   EURAXESS</a> ( <a href="https://euraxess.ec.europa.eu/career-development/researchers/manual-scientific-entrepreneurship/major-steps/trl">https://euraxess.ec.europa.eu/career-development/researchers/manual-scientific-entrepreneurship/major-steps/trl</a> )

### Guidance on the use of generative AI tools for the preparation of the proposal

When considering the use of generative artificial intelligence (AI) tools for the preparation of the proposal, it is imperative to exercise caution and careful consideration. The AI-generated content should be thoroughly reviewed and validated by the applicants to ensure its appropriateness and accuracy, as well as its compliance with intellectual property regulations. Applicants are fully responsible for the content of the proposal (even those parts produced by the AI tool) and must be transparent in disclosing which AI tools were used and how they were utilized.

Specifically, applicants are required to:

- Verify the accuracy, validity, and appropriateness of the content and any citations generated by the AI tool and correct any errors or inconsistencies.
- Provide a list of sources used to generate content and citations, including those generated by the AI tool. Double-check citations to ensure they are accurate and properly referenced.
- Be conscious of the potential for plagiarism where the AI tool may have reproduced substantial text from other sources. Check the original sources to be sure you are not plagiarizing someone else's work.
- Acknowledge the limitations of the AI tool in the proposal preparation, including the potential for bias, errors, and gaps in knowledge.

**⚠️ Fill in the title of your proposal below.**

<b>TITLE OF THE PROPOSAL</b>
------------------------------

[This document is tagged. Do not delete the tags; they are needed for processing.] #@APP-FORM-HECSA@#

**List of participants**

Participant	Participant organisation name	Country
1 (Coordinator)		
2 Affiliated entity(ies), if any. Please explain the link with the coordinator and the role in the project.		

[This document is tagged. Do not delete the tags; they are needed for processing.] #@APP-FORM-HEEICTRA@#

**1. Excellence** #@REL-EVA-RE@#

***Excellence – aspects to be taken into account***

- **Technological breakthrough:** Does the technology have a high degree of novelty and higher performance compared to other technologies available or in development? Does the technology present high commercial potential?
- **Objectives:** How credible and feasible are the objectives for the planned technology development and maturation? How credible and feasible are the objectives and KPIs for the planned business development process?
- **Methodology:** Is the timing right for this technology/innovation (i.e., feasibility, technological readiness level, (TRL)?

**1.1 Technological breakthrough** #@PRJ-OBJ-PO@#

- a. Describe the degree of novelty and performance compared to other technologies available or in development, and the competitive advantages. . In other words, how is it done today and what is the limit of the current practice?<sup>1</sup>
- b. Describe the breakthrough nature of the innovation in this proposal. What is new in your approach and why do you think it will be successful?
- c. Describe the commercial potential of the technology.

**1.2 Objectives**

- a. Describe the objectives both for the technology development and for the business validation and development of the project. Describe what you are trying to do. Articulate your objectives using very little

<sup>1</sup> <https://www.darpa.mil/about/heilmeier-catechism>

or preferably no jargon. Objectives should be specific, measurable, achievable, relevant and time bound within the duration of the project:

- Describe the KPIs for measuring the achievement of technology development objectives.
- Describe the KPIs for measuring the achievement of the business validation and development objectives.
- How appropriate are the objectives for the planned technology development and validation of the innovation in relevant application environments?
- Describe how the specific operational objectives address the challenges and unknown to bring the innovation to the public.

**1.3 Methodology** #@CON-MET-CM@# #@COM-PLE-CP@#

- a. Describe and explain the approach and methodology including the concepts, models and assumptions that will enable you to deliver your project’s objectives. Explain why they are well-suited and come at the right time to handle the significant unknowns and uncertainties related to technology and innovation developed in the project. Explain how appropriate they are to enable alternative directions and options if needed (e.g. in case certain risk materialize).

**⚠** *This section should be presented as a narrative. The detailed tasks and work packages are described below under ‘Implementation’.*

Elaborate and substantiate on the current TRL of your innovation at system level AND in relation to the intended final user application (e.g. clinical for for health innovation, consumer for energy systems, etc.).

Describe the milestones already achieved in the development of the technology.

Elaborate on the final TRL of the innovation aimed for at the end of the project.

What is you Unique Selling Point and how do you intend to further develop it?

**⚠** *Where relevant, include how the project methodology complies with the ‘do no significant harm’ principle as per Article 17 of [Regulation \(EU\) No 2020/852](#) on the establishment of a framework to facilitate sustainable investment (i.e. the so-called ‘EU Taxonomy Regulation’). This means that the methodology is designed in a way it is not significantly harming any of the six environmental objectives of the EU Taxonomy Regulation.*

⚠ *If you plan to use, develop and/or deploy artificial intelligence (AI) based systems and/or techniques you must demonstrate their technical robustness. AI-based systems or techniques should be, or be developed to become:*

- *technically robust, accurate and reproducible, and able to deal with and inform about possible failures, inaccuracies and errors, proportionate to the assessed risk they pose*
- *socially robust, in that they duly consider the context and environment in which they operate*
- *reliable and function as intended, minimizing unintentional and unexpected harm, preventing unacceptable harm and safeguarding the physical and mental integrity of humans*
- *able to provide a suitable explanation of their decision-making processes, whenever they can have a significant impact on people's lives.*

b. Briefly describe how the gender dimension (i.e. sex and/or gender analysis) is taken into account in the project's research and innovation content . If you do not consider such a gender dimension to be relevant in your project, please provide a brief justification.

⚠ *This question relates to the content of the planned research and innovation activities, and not to gender balance in the teams in charge of carrying out the project.*

⚠ *Sex and gender analysis refers to biological characteristics and social/cultural factors respectively. For guidance on methods of sex / gender analysis and the issues to be taken into account, please refer to <https://op.europa.eu/en/publication-detail/-/publication/33b4c99f-2e66-11eb-b27b-01aa75ed71a1/language-en>*

c. Briefly describe how appropriate open science practices are implemented as an integral part of the proposed methodology. Show how the choice of practices and their implementation are adapted to the nature of your work, in a way that will increase the chances of the project delivering on its objectives.

If you believe that none of these practices are appropriate for your project, please provide a brief justification here.

⚠ *Open Science is an approach based on open cooperative work and systematic sharing of knowledge and tools as early and widely as possible in the process. Open science practices include early and open sharing of research (for example through preregistration, registered reports, pre-prints, or crowd-sourcing); research output management; measures to ensure reproducibility of research outputs; providing open access to research outputs (such as publications, data, software, models, algorithms, and workflows); participation in open peer-review; and involving all relevant knowledge actors including citizens, civil society and end users in the co-creation of R&I agendas and contents (such as citizen science).*

⚠ *Please note that this question does not refer to outreach actions that may be planned as part of communication, dissemination and exploitation activities. These aspects should instead be described below under 'Impact'.*

d. Research data management and management of other research outputs: Applicants generating/collecting data and/or other research outputs (except for publications) during the project must provide a very short description on how the data/research outputs will be managed:

**Types of data/research outputs** (e.g. experimental, observational, images, text, numerical) and their estimated size; if applicable, combination with, and provenance of, existing data.

**Findability of data/research outputs:** Types of persistent and unique identifiers (e.g. digital object identifiers) and trusted repositories that will be used.

**Accessibility of data/research outputs:** IPR considerations and timeline for open access (if open access not provided, explain why); provisions for access to restricted data for verification purposes.

**Interoperability of data/research outputs:** Standards, formats and vocabularies for data and metadata.

**Reusability of data/research outputs:** Licenses for data sharing and re-use (e.g. Creative Commons, Open Data Commons); availability of tools/software/models for data generation and validation/interpretation /re-use.

**Curation and storage/preservation costs;** person/team responsible for data management and quality assurance.

**⚠️** *Proposals selected for funding under Horizon Europe will need to develop a detailed data management plan (DMP) for making their data/research outputs findable, accessible, interoperable and reusable (FAIR) as a deliverable by month 6 and revised towards the end of a project’s lifetime. It is allowed not to publish any data, if this might harm the protection of the results or commercialisation strategy.*

**⚠️** *For guidance on Open Science practices and research data management, please refer to the relevant section in the [EIC work programme](#) and in the [HE Programme Guide on the Funding & Tenders Portal](#).*

#\$PRJ-OBJ-POS# #SREL-EVA-RE\$# #@CON-MET-CM@# #@COM-PL-CP@#

**2. Impact** #@IMP-ACT-IA@#

**Impact – aspects to be taken into account**

- Credibility of the impacts: To what extent the commercial impact(s) described in the proposal are credible and substantial within the project and beyond (e.g., one or several sectors, setting new standards, etc.)?
- Economic and/or societal benefits: To what extent does the proposed innovation have scale-up potential including high capacity to gain or create new markets? To what extent is the proposed innovation expected to generate positive impacts for Ukraine (e.g., strategic autonomy, employment etc.)?

**⚠️** *In this section you should show how your project could contribute to the outcomes and impacts described in the work programme, the likely scale and significance of this contribution, and the measures to maximise these impacts.*

**2.1 Credibility of the impacts**

- Describe how the activities proposed support the development of a business model and the product features. In particular describe the methodology to validate the initial/incipient business model.

- Describe how the activities support the commercialisation and other relevant aspects (intellectual property rights, regulation, certification and standardisation).
  - Outline your strategy for the management of intellectual property, foreseen protection measures, such as patents, design rights, copyright, trade secrets etc., and how these would be used to support exploitation.
  - Describe how you performed early exploration of potential markets for your innovation to test potential demand and acceptability, as well as potential competitors. Provide or quote any support data you may have.
  - Describe how you will validate the problem / solution fit.
  - Describe the planned measures to maximise the impact of your project by providing a brief version of your 'plan for the dissemination and exploitation including communication activities'. Briefly describe the dissemination, exploitation and communication measures that are planned, and the target group(s) addressed (e.g. scientific community, end users or potential customers, financial actors, public at large).
- ⚠** *In case your proposal is selected for funding, a more detailed 'plan for dissemination and exploitation including communication activities' will need to be provided as a mandatory project deliverable within 6 months after the start of the project. This plan shall be periodically updated in alignment with the project's progress.*
- ⚠** *Communication<sup>2</sup> measures should promote the project throughout the full lifespan of the project. The aim is to inform and reach out to society and show the activities performed, and the use and the benefits the project will have for citizens. Activities must be strategically planned, with clear objectives, start at the outset and continue through the lifetime of the project. The description of the communication activities needs to state the main messages as well as the tools and channels that will be used to reach out to each of the chosen target groups.*
- ⚠** *All measures should be proportionate to the scale of the project, and should contain concrete actions to be implemented both during and after the end of the project. In the justification, explain why each measure chosen is best suited to reach the target group addressed.*
- ⚠** *If exploitation is expected primarily in non-associated third countries, justify why and explain how that exploitation is still in the Union's common interest.*
- ⚠** *If applicable, describe possible feedback to policy measures generated by the project that will contribute to designing, monitoring, reviewing and rectifying (if necessary) existing policy and programmatic measures or shaping and supporting the implementation of new policy initiatives and decisions.*

## 2.2 Economic and/or societal benefits

- Describe how the proposed innovation and its related activities can have scale up potential to gain and create new European or global markets. Be concrete and realist providing arguments and data to support your assertions. In other words, if you are successful what difference will your innovation/technology make and to whom?
- Describe concretely the positive impacts for the European Union, Member States or Associated Countries

---

<sup>2</sup> For further guidance on communicating EU research and innovation for project participants, please refer to the [Online Manual](#) on the Funding & Tenders Portal.

(e.g. strategic autonomy, productivity gains, green deal, etc.). If possible, provide data to support your claims.

**2.3 Investment readiness**

- Please describe the proposed measures to become investment ready and develop plans to commercialise the project outcomes (including through IP management).
- Please describe how you intend to validate your value proposition with potential users and customers. Please describe the go-to-market pathway/strategy, what regulatory approvals might impact your go-to-market strategy, time to market, possible business and revenue model. *Please select in the box below the go-to-market pathway.*
- *If you plan to raise private capital during or shortly after the project end, please mention any concrete activities you plan to execute together with any preparation (if) needed for a successful outcome.*

<p>a. Describe the path to market of the innovation beyond the duration of the project</p>
<p><i>The coordinator will directly exploit the innovation. Elaborate.</i></p> <p><i>or</i></p> <p><i>The coordinator will create a new company. Elaborate and be precise. who, how, when.</i></p> <p><i>or</i></p> <p><i>The coordinator will licence the technology to an established company. Elaborate.</i></p> <p><i>or</i></p> <p><i>None of the above or a combination of the above. Elaborate the different alternative and be as precise and concrete as possible.</i></p>

#§IMP-ACT-IA§#

**3. Quality and efficiency of the implementation #@QUA-LIT-QL@# #@CON-SOR-CS@# #@PRJ-MGT-PM@#**

<p><b>Quality and efficiency of the implementation – aspects to be taken into account</b></p> <ul style="list-style-type: none"> <li>– <u>Quality and motivation of the team</u>: To what extent does the team have the necessary high-quality capabilities and high motivation to move decisively towards market. To what extent does the applicant team have the necessary expertise to create a unique commercial value from the emerging technology and develop an attractive business and investment proposition?</li> <li>– <u>KPIs and Milestones</u>: Are both milestones and KPIs present, relevant and clearly defined (measurable, timed, comparable etc.) to track progress along the pathway towards objectives? Have the main risks (e.g., technological, market, financial etc.) been identified, together with measures to mitigate them, in order to achieve the project objectives?</li> <li>– <u>Workplan and allocation of resources</u>: How appropriate and effective is the allocation of resources (person-months and equipment) in the workplan and work packages?</li> </ul>
--

**Patent families** (please list here all granted patents / ongoing patent applications that may be relevant for the evaluation)

Patent Applicant	Number	Title of patent	Date of filing/Priority date

### 3.1 Quality and motivation of the team

- Explain to what extent does(do) the applicant(s) bring the necessary high-quality expertise, capabilities and motivation to create a unique commercial value from the emerging technology and develop an attractive business and investment proposition. Include in the description affiliated entities and associated partners, if any.
- If applicable explain, for the main exploitation partner who is and what experience have:
  - The CEO
  - The CSO, chief scientific officer
  - The CFO, chief financial officer
  - The rest of the main team members and their unique expertise.
- If not (yet) applicable explain how you intend to acquire them or why they are not needed.

### 3.2 KPIs, milestones and risks

- Describe technological (e.g., performance) and go-to market milestones. Please include here also KPIs necessary to measure the achievement of the milestones (please include a narrative in this section and summarise it in table 3.3d). Include for example number of meetings with potential customers to validate initial business plan or other means of interacting with potential customers, market experts, investors and potential partners to mature your business model via market validation. Maturing both the technology and business model is a key characteristic and requirement of this call!
- Describe the critical technical and market-related risks, which are relevant to track progress along the pathway towards objectives. Detail any risk mitigation measures (please include a short narrative in this section and summarise it in table 3.3e). Please include here any KPI necessary to measure the achievement of the milestones.

### 3.3 Work plan and resources #@WRK-PLA-WP@#

- Please describe the work plan (work packages, tasks, deliverables, time-line, etc.):
  - brief presentation of the overall structure of the work plan;
  - timing of the different work packages and their components (Gantt chart or similar);

**⚠ Please use the below table when planning Reporting Periods for your project:**

Project duration	Number of periods	RP1 duration	RP2 duration	RP3 duration	RP4 duration
12	1	12	-	-	-
18	1	18	-	-	-
24	2	12	12	-	-
30	2	12	18	-	-
36	2	12	24	-	-

(after 36 months, reporting occurs every 12 months.)

- graphical presentation of the components showing how they inter-relate (Pert chart or similar);
- detailed work description, i.e.:
  - a list of work packages (table 3.3a);
  - a description of each work package (table 3.3b);
  - a list of deliverables (table 3.3c);
- *Give full details. Base your account on the logical structure of the project and the stages in which it is to be carried out. Each work package should be a substantial part of the work plan, and the number of work packages should be proportionate to the scale and complexity of the project.*
- *Structure each work package by breaking it down into tasks. If tasks are not appropriate, work packages can be organised according to other criteria (e.g., according to the type of work or thematically). For each task or element of the work package, describe all activities to be carried out and quantify them (e.g., number of protocols, tests, measurements, combinations, study subjects, conferences, publications, etc.). Provide enough detail to clarify who will do this work and why it is needed for the project, (e.g., the level of qualification and number of person-months for personnel, as well as the requested equipment, consumables, meetings, etc.), to justify the proposed resources and so that progress can be monitored, including by the Commission.*
- *Resources (person-months) assigned to work packages should be in line with their objectives and deliverables. You are advised to include a distinct work package on 'project management', and to give due visibility in the work plan to 'data management' 'dissemination and exploitation' and 'communication activities', either with distinct tasks or distinct work packages.*
- *You will be required to update the 'plan for the dissemination and exploitation of results including communication activities', and a 'data management plan', (this does not apply to topics where a plan was not required.) This should include a record of activities related to dissemination and exploitation that have been undertaken and those still planned.*
- *Please make sure the information in this section matches the costs as stated in the budget table in section 3 of the application forms, and the number of person months, shown in the detailed work package descriptions.*
- a list of Key Performance Indicators (KPI) and milestones (table 3.3d);

- a list of critical risks, relating to project implementation, that the stated project's objectives may not be achieved. Detail any risk mitigation measures. You will be able to update the list of critical risks and mitigation measures as the project progresses (table 3.3e);
- a table showing number of person months required (table 3.3f);
- a table showing description and justification of subcontracting costs for each participant (table 3.3g);
- a table showing justifications for 'purchase costs' (table 3.3h) for participants where those costs exceed 15% of the personnel costs (according to the budget table in the proposal);
- if applicable, a table showing justifications for 'other costs categories' (table 3.3i);
- if applicable, a table showing in-kind contributions from third parties (table 3.3j).
- If you want to deviate from the maximum recommended budget (Euro 2.5 mln) and/ or duration (36 months), please DO explain AND motivate why this is really necessary.

#§CON-SOR-CS§# #§PRJ-MGT-PMS#

Example, not to be completed

**Tables for section 3.3**

**⚠** Use plain text for the tables in section 3.3. If the proposal is invited to start Grant Agreement preparation, these tables will have to be encoded in the grant management IT tool, where no graphics or special formats are supported.

**Table 3.3a: List of work packages**

*In lump sums, payment can be claimed after the completion of a work package. Therefore, should you wish to get a payment after the first reporting period (month 12), please ensure you complete a work package at month 12."*

Work package No	Work Package Title	Lead Participant No	Lead Participant Short Name	First name and family name of the Work Package Leader	Gender of Work Package Leader	Person-Months	Start Month	End month
						Total person-months		

Example, not to be completed

**Table 3.3b: Work package description**

For each work package:

<b>Work package number</b>	
<b>Work package title</b>	

⚠ *Participants involved in each WP and their efforts are shown in table 3.3f. Lead participant and starting and end date of each WP are shown in table 3.3a.)*

**Objectives**

**Description of work** (where appropriate, broken down into tasks), lead partner and role of participants. For each task, quantify the amount of work. Provide enough detail to justify the resources requested and clarify why the work is needed and who will do it. Deliverables linked to each WP are listed in table 3.3c (no need to repeat the information here).

Example, not to be completed

**Table 3.3c: List of Deliverables**

Only include deliverables that meaningful to the intended outcome of the project (e.g. no project hand book). The content of the technical deliverables should target scientific or technology expert monitors and focus on outcomes and results rather than description of the activities and processes.

You must include as distinct deliverables

- data management plan (DMP) by month 6<sup>3</sup>,
- Draft business plan (including a draft Business Model Canvas) by month 12
- Final business plan (including a draft Business Model Canvas) by the end of the project
- Report on IP management (ideally a FTO analysis to be performed during the execution of the project)
- Plan for dissemination and communication activities within the first 6 months of the project.

Number	Deliverable name	Short description	Work package number	Short name of lead participant	Type	Dissemination level	Delivery date (in months)

**KEY**

*Deliverable numbers in order of delivery dates. Please use the numbering convention <WP number>.<number of deliverable within that WP>. For example, deliverable 4.2 would be the second deliverable from work package 4.*

**Type:**

*Use one of the following codes:*

- R: Document, report (excluding the periodic and final reports)
- DEM: Demonstrator, pilot, prototype, plan designs
- DEC: Websites, patents filing, press & media actions, videos, etc.
- DATA: Data sets, microdata, etc.
- DMP: Data management plan
- ETHICS: Deliverables related to ethics issues.

<sup>3</sup> *Proposals selected for funding under Horizon Europe will need to develop a detailed data management plan (DMP) for making their data/research outputs findable, accessible, interoperable and reusable (FAIR) as a deliverable by month 6 and revised towards the end of a project’s lifetime. It is allowed not to publish any data, if this might harm the protection of the results or commercialisation strategy. The DMP will evolve during the lifetime of the project in order to present the status of the project’s reflections on data management. A template for such a plan is available in the Online Manual on the Funding & Tenders Portal.*

**SECURITY:** Deliverables related to security issues  
**OTHER:** Software, technical diagram, algorithms, models, etc.

**Dissemination level:**

Use one of the following codes:

PU – Public, fully open, e.g. web (Deliverables flagged as public will be automatically published in CORDIS project’s page)

SEN – Sensitive, limited under the conditions of the Grant Agreement

Classified R-UE/EU-R – EU RESTRICTED under the Commission Decision No2015/444

Classified C-UE/EU-C – EU CONFIDENTIAL under the Commission Decision No2015/444

Classified S-UE/EU-S – EU SECRET under the Commission Decision No2015/444

**Delivery date**

Measured in months from the project start date (month 1)

**Table 3.3d: List of milestones**

Milestone number	Milestone name	Related work package(s)	Due date (in month)	Means of verification (KPI)

Make sure regular and meaningful milestones are set every six months (see also the definition of the milestones at the beginning of the template). Milestones must be major achievements, decision points on whether the work (objectives and tasks) should continue as planned, modified or terminated. All milestones should be associated with deliverables and/or additionally be phrased in specific, quantitative terms where possible and relevant (means of verification (KPI)), which will verify the attainment of the milestones. In other words, how long will it take and what are the intermediate and final checks/verifications for success?

**KEY**

**Due date**

Measured in months from the project start date (month 1)

**Means of verification**

Show how you will confirm that the milestone has been attained. Refer to indicators if appropriate. For example: a laboratory prototype that is ‘up and running’; software released and validated by a user group; field survey complete and data quality validated. Please include here any KPI necessary to measure the achievement of the milestones.

**Table 3.3e: Critical risks for implementation #@RSK-MGT-RM@#**

Description of risk (indicate level of (i) likelihood, and (ii) severity: Low/Medium/High)	Work package(s) involved	Proposed risk-mitigation measures

**Definition critical risk:**  
*A critical risk is a plausible event or issue that could have a high adverse impact on the ability of the project to achieve its objectives.*

**Level of likelihood to occur: Low/medium/high**  
*The likelihood is the estimated probability that the risk will materialise even after taking account of the mitigating measures put in place.*

**Level of severity: Low/medium/high**  
*The relative seriousness of the risk and the significance of its effect.*

#§RSK-MGT-RM§#

**Table 3.3f: Summary of staff effort**

Please indicate the number of person/months over the whole duration of the planned work, for each work package, for each participant. Identify the work-package leader for each WP by showing the relevant person-month figure in bold.

	WPn	WPn+1	WPn+2	Total Person-Months per Participant
Participant Number/Short Name				
ParticipantNumber/Short Name				
Participant Number/Short Name				
<b>Total Person Months</b>				

**Table 3.3g: ‘Subcontracting costs’ items**

For each participant describe and justify the tasks to be subcontracted (please note that core tasks of the project should not be sub-contracted).

Participant Number/Short Name		
	Cost (€)	Description of tasks and justification
<b>Subcontracting</b>		

**Table 3.3h: ‘Purchase costs’ items (travel and subsistence, equipment and other goods, works and services)**

Please complete the table below for each participant if the purchase costs (i.e. the sum of the costs for ‘travel and subsistence’, ‘equipment’, and ‘other goods, works and services’) exceeds 15% of the personnel costs for that participant (according to the budget table in the proposal part). The record must list cost items in order of costs and starting with the largest cost item, up to the level that the remaining costs are below 15% of personnel costs.

Please be aware that CFS and audit costs are not needed and therefore not eligible in lump sum.

Participant Number/Short Name		
	Cost (€)	Justification
Travel and subsistence		
Equipment		
Other goods, works and services		
Remaining purchase costs (<15% of pers. Costs)		
<b>Total</b>		

**Table 3.3i: ‘Other costs categories’ items (e.g. internally invoiced goods and services)**

Please complete the table below for each participant that would like to declare costs under other costs categories (e.g. internally invoiced goods and services), irrespective of the percentage of personnel costs.

Participant Number/Short Name		
	Cost (€)	Justification
Internally invoiced goods and services		
...		

**Table 3.3j: ‘In-kind contributions’ provided by third parties**

Please complete the table below for each participant that will make use of in-kind contributions (non-financial resources made available free of charge by third parties). In kind contributions provided by third parties free of charge are declared by the participants as eligible direct costs in the corresponding cost category (e.g. personnel costs or purchase costs for equipment).

Participant Number/Short Name			
Third party name	Category	Cost (€)	Justification
	<b>Select between</b> Seconded personnel Travel and subsistence Equipment Other goods, works and services Internally invoiced goods and services		

#SQUA-LIT-QLS# #SWRK-PLA-WPS#

## ANNEXES TO PROPOSAL PART B

The annexes to be uploaded for this call are (standard templates are published in the Funding & Tenders portal):

- **LUMP SUM DETAILED BUDGET TABLE**
- **CLINICAL TRIALS:** Annex with information on clinical trials
- **ETHICS:** ethics self-assessment should be included in proposal part A. However, in calls where several serious ethics issues are expected, the character limited in this section of proposal part A may not be sufficient for participants to give all necessary information. In those cases, participants may include additional information in an annex to proposal part B.

Example, not to be completed