

The EU Framework Programme for Research and Innovation



# Marie Skłodowska-Curie Actions Individual Fellowships (IF) – European fellowships (EF)

# Administrative forms (Part A) Research proposal (Part B)

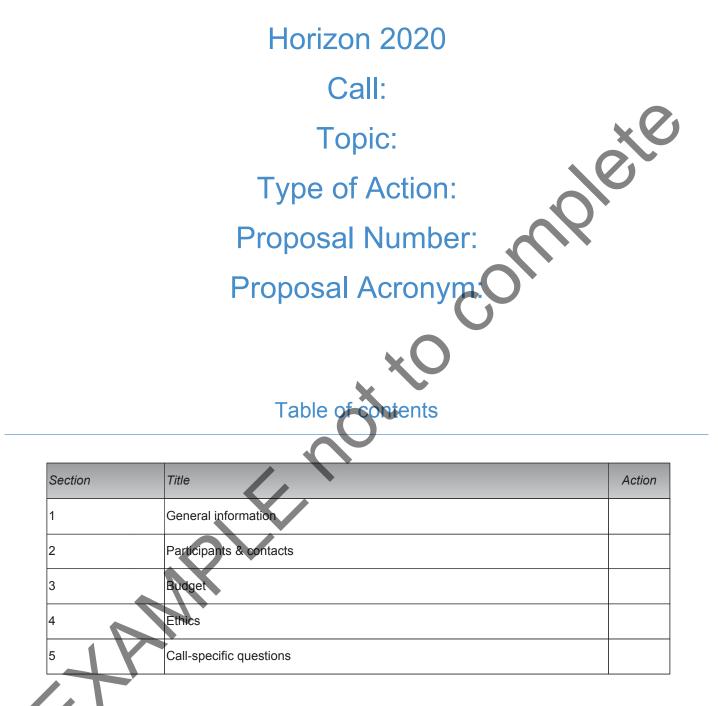
Version 2.0 30 March 2015

#### Disclaimer

This document is aimed at informing potential applicants for Horizon 2020 funding. It serves only as an example. The actual Web forms and templates, provided in the online proposal submission system under the Participant Portal, might differ from this example. Proposals must be prepared and submitted .via the online proposal submission system under the Participant Portal.







### How to fill in the forms?

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

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1 0 0 1	aral information		

#### 1 - General information

Торіс		Type of action
Call identifier		Acronym
Proposal title	The title should be no lo the non-specialist in yo	onger than 200 characters (with spaces) and should be understandable to ur field.
	ote that for technical rea e removed: < > " &	isons, the following characters are not accepted in the Proposal Title and will
	Duration in months	
Panel		
Please select up to 5 of relevance.	descriptors (and at leas	t 1) that best characterise the subject of your proposal, in descending order
Descriptor 1		Add
Free keywords		er of keywords that you consider necessary to characterise the scope of a limit of 200 characters.
Abstract		
<ul> <li>the objective</li> <li>how they wil</li> <li>their relevan</li> <li>Will be used as the s</li> <li>management commi</li> <li>Do not inclus</li> <li>Use plain typ</li> </ul>	es of the proposal I be achieved ce to the work program short description of the p ttees and other intereste de any confidential infor ped text, avoiding formu	roposal in the evaluation process and in communications with the programme ed parties .
Remaining character	s 2000	
Has this proposal (or	a very similar one) bee	n submitted in the past 2 years in response to a call for

proposals under the 7th Framework Programme, Horizon 2020 or any other EU programme(s)? C Yes C No

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	osal (or a very similar one) been submitted to a Horizon 2020 Marie Skłodowska- al Fellowship call?	○ Yes ○No
	AMPLE NOT	

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#### Declarations

1) The applicant (future beneficiary) declares to have the explicit consent of all partner organisations (if applicable) on their participation and on the content of this proposal.

2) The information contained in this proposal is correct and complete.

3) This proposal complies with ethical principles (including the highest standards of research integrity — as set out, for instance, in the European Code of Conduct for Research Integrity — and including, in particular, avoiding fabrication, falsification, plagiarism or other research misconduct).

4) The applicant (future beneficiary) confirms:

, the first of the	
- to have carried out the self-check of the financial capacity of the organisation on https://ec.europa.eu/research/participants/portal/desktop/en/organisations/lfv.html or to be covered by a financial viability check in an EU project for the last closed financial year. Where the result was "weak" or "insufficient", the applicant (future beneficiary) confirms being aware of the measures that may be imposed in accordance with the H2020 Grants Manual (Chapter on Financial capacity check); or	0
- is exempt from the financial capacity check being a public body including international organisations, higher or secondary education establishment or a legal entity, whose viability is guaranteed by a Member State or associated country, as defined in the H2020 Grants Manual (Chapter on Financial capacity check); or	0
- as sole participant in the proposal is exempt from the financial capacity check.	0

5) The applicant (future beneficiary) hereby declares:

- it is fully eligible in accordance with the criteria set out in the specific call for proposals; and	
- it has the financial and operational capacity to carry out the proposed action.	

The applicant (future beneficiary) is only responsible for the correctness of the information relating to his/her own organisation. Where the proposal to be retained for EU funding, the applicant (future beneficiary) will be required to present a formal declaration in this respect.

According to Article 131 of the Financial Regulation of 25 October 2012 on the financial rules applicable to the general budget of the Union (Official Journal L 298 of 26.10.2012, p. 1) and Article 145 of its Rules of Application (Official Journal L 362, 31.12.2012, p. 1) applicants found guilty of misrepresentation may be subject to administrative and financial penalties under certain conditions.

#### Personal data protection

Your reply to the grant application will involve the recording and processing of personal data (such as your name, address and CV), which will be processed pursuant to Regulation (EC) No 45/2001 on the protection of individuals with regard to the processing of personal data by the Community institutions and bodies and on the free movement of such data. Unless indicated otherwise, your replies to the questions in this form and any personal data requested are required to assess your grant application in accordance with the specifications of the call for proposals and will be processed solely for that purpose. Details concerning the processing of your personal data are available on the privacy statement. Applicants may lodge a complaint about the processing of their personal data with the European Data Protection Supervisor at any time.

Your personal data may be registered in the Early Warning System (EWS) only or both in the EWS and Central Exclusion Database (CED) by the Accounting Officer of the Commission, should you be in one of the situations mentioned in: -the Commission Decision 2008/969 of 16.12.2008 on the Early Warning System (for more information see the <u>Privacy Statement</u>), or -the Commission Regulation 2008/1302 of 17.12.2008 on the Central Exclusion Database

(for more information see the <u>Privacy Statement</u>).

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### List of participants

#	Participant Legal Name	Country
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2 - Administrative data of participating organisations			

### **Future Host Institution**

PIC	Legal name		ר
Short name:			
Address of the	organisation		
Stree	et		$\sim$
Tow	n		
Postcod	e		•
Country	y	U	
Webpage	e	×O	
Legal Status o	f your organisation		
		<b>N</b>	
Research and I	nnovation legal statuses	$\sim$	
Public body	no	<ul> <li>Legal person</li> </ul>	no
Non-profit	ho		
International organ	nisationno		
	nisation of European interest no		
	ner education establishment no		
	ationno		
Small and Mediun Academic Sector	n-sized Enterprises (SMEs) no		
Nace code			

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#### Department(s) carrying out the proposed work

#### Department 1

Department name	×0
	Same as organisation address
Street	Please enter street name and number.
Town	
Postcode	
Country	
although the propos	Department carrying out the proposed work is not the same as the location of the Host Institute, please note that al submission system calculates the budget of the project based on the location of the Host Institute, the budget of ant agreement will be calculated by using the country coefficient of the location of the Department carrying out the

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#### Supervisor

The name and e-mail of the Researcher and Supervisor are read-only in the administrative form, only additional details can be edited here. To give access rights and contact details of contact persons, please go back to Step 4 of the submission wizard and save the changes.

Title	Sex O Male O Fe	emale
First name*	Last name*	
E-Mail*		
Position in org.	Please indicate the position of the Contact Point above in the organisation	K
Department	Please indicate the department of the Contact Point above in the organisati	
	□ Same as organisation address	
Street		
Town	Post code	
Country		
Website		
Phone	+xxx xxxxxxx Phone 2 +xxx xxxxxxx Fax +xxx	XXXXXXXXX

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R	е	S	е	а	r	C	h	е	r

The name and e-mail of the Researcher and Supervisor are read-only in the administrative form, only additional details can be edited here. To give access rights and contact details of contact persons, please go back to Step 4 of the submission wizard and save the changes.

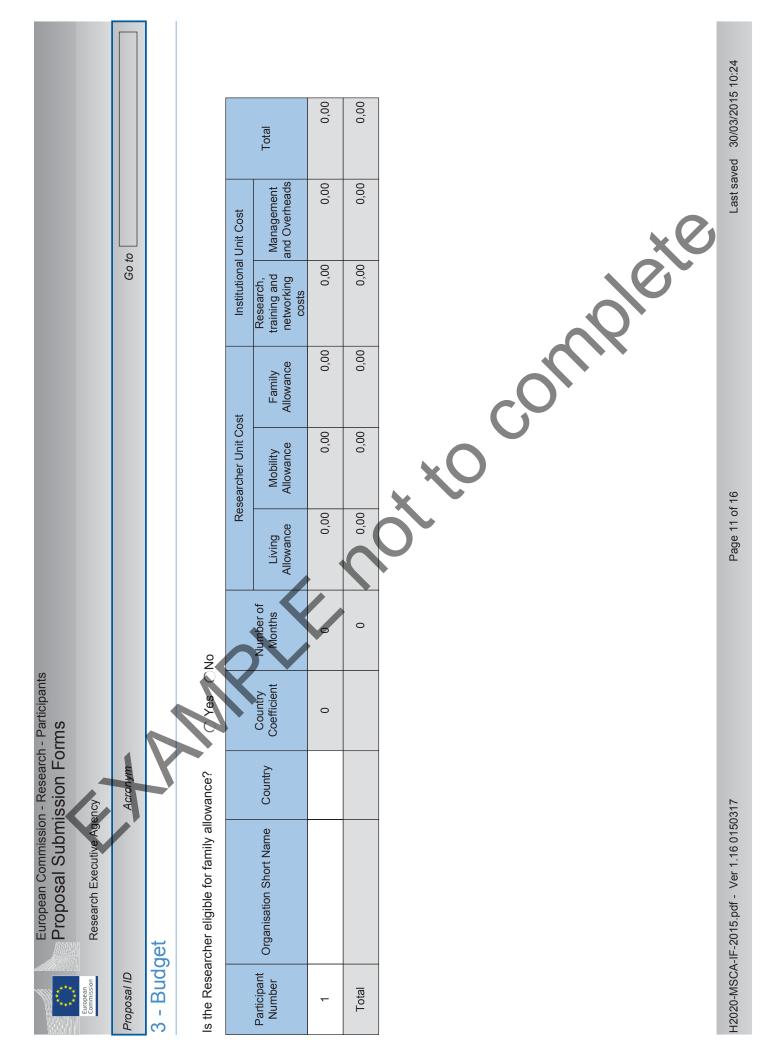
Researcher ID	If you have a researcher identifier num	nber (e.g. ResearcherID, ORC	ID) please enter it here.
Last Name*		Last Name at Birth	
First Name(s)*		Gender*	∩ Male ⊂ Female
Title		Country of residence	ce*
Nationality*		Nationality 2	
Date of Birth (DD/	MM/YYYY)	Country of Birth*	
		Place of Birth	
Contact addre	SS	×O	Same as organisation address
		<u> </u>	
Current organisa	ition name		
Current Departm Laboratory name	nent/Faculty/Institute/		
Street	Please enter street name and n	umber.	
Postcode/Cedex		Town	
Phone	+XXX XXXXXXXXX	Country	
Phone2 / Mobile			
E-Mail*	P1.		
Qualifications			
University Degre	e	Date of award (D	D/MM/YYYY)
Doctorate (in pro	gress)	Date of award (D	D/MM/YYYY)
Doctorate		Date of award (D	D/MM/YYYY)
Full time postgra	duate research experience	Number of mor	nths
Other Academic	qualifications	Date of award (D	DD/MM/YYYY)

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#### Place of activity/place of residence (previous 5 years - most recent one first)

Indicate the period(s) and the country/contries in which you have legally resided and/or had your main activity (work, status, ..) during the last 5 years up until the deadline for the submission of the proposal. Please fill in this section without gaps, until the call deadline (11/09/2014).

Period from	Period to	Duration (days)	Country	Add
	11/09/2014			Remove
		Total		0
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Short name null \* Proposal \* Ethics \* ethics\_title \* sub

#### 4 - Ethics issues table

1. HUMAN EMBRYOS/FOETUSES		Page
Does your research involve Human Embryonic Stem Cells (hESCs)?	⊖Yes	. 0.
Does your research involve the use of human embryos?	⊖Yes ●No	X
Does your research involve the use of human foetal tissues / cells?	⊖Yes ⊙No	0
2. HUMANS		Page
Does your research involve human participants?	⊖Yes ⊙No	
Does your research involve physical interventions on the study participants?	OYes ⊙No	
3. HUMAN CELLS / TISSUES		Page
Does your research involve human cells or tissues (other than from Human Embryos/ Foetuses, i.e. section 1)?	⊖Yes ⊙No	
4. <u>PERSONAL DATA</u> (ii)		Page
Does your research involve personal data collection and/or processing?	⊖Yes ●No	
Does your research involve further processing of previously collected personal data (secondary use)?	⊖Yes	
5. <u>ANIMALS</u> (iii)		Page
Does your research involve animals?	⊖Yes ⊙No	

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6. THIRD COUNTRIES			Page
Does your research involve non-EU countries?	⊖ Yes	No	
Do you plan to use local resources (e.g. animal and/or human tissue samples, genet material, live animals, human remains, materials of historical value, endangered fauna flora samples, etc.)? (v)		No	XO
Do you plan to import any material from non-EU countries into the EU? For data imports, please fill in also section 4. For imports concerning human cells or tissues, fill in also section 3.	⊖Yes	© No	Ø
Do you plan to export any material from the EU to non-EU countries? For data exports, please fill in also section 4. For exports concerning human cells or tissues, fill in also section 3.	Yes	• No	
If your research involves low and/or lower middle income countries, are benefits-sharing measures foreseen? (vii)	OYes	No	
Could the situation in the country put the individuals taking part in the research at risk?	⊖Yes	No	
7. ENVIRONMENT & HEALTH and SAFETY See legal references at the end of the section. (vi)			Page
Does your research involve the use of elements that may cause harm to the environment, to animals or plants? For research involving animal experiments, please fill in also section 5.	ne 🔿 Yes	No	
Does your research deal with endangered fauna and/or flora and/or protected areas?	⊖ Yes	No	
Does your research involve the use of elements that may cause harm to human including research staff? For research involving human participants, please fill in also section 2.	s, OYes	No	
8. <u>DUAL USE</u> (vii)			Page
Does your research have the potential for military applications?	⊖ Yes	No	
9. MISUSE			Page
Does your research have the potential for malevolent/criminal/terrorist abuse?	⊖ Yes	No	
10. OTHER ETHICS ISSUES			Page
Are there any other ethics issues that should be taken into consideration? Please specif	<sup>fy</sup> O Yes	No	

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I confirm that I have taken into account all ethics issues described above and that, if any ethics issues apply, I will complete the ethics self-assessment and attach the required documents.

How to Complete your Ethics Self-Assessment

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5 - Call specific questions	
Eligibility Researcher (future fellow)	
1. Were you in the last 5 years in military service?	⊖Yes ⊖No
2. Are you a national of Member State or Associated Country?	⊖ Yes ⊖No
Other Questions	$\mathbf{V}$
For communication purposes only, the REA asks for permission to publish the name of the researce the proposal be retained for funding.	cher (future fellow) should
1. Does the researcher (future fellow) give this permission?	○Yes ○No
2. Is there a secondment in Member States or Associated Countries evisaged in Part B of this proposal?	○ Yes ○No
Attention: this secondment is different than the outgoing phase in the Third Country, and only takes place an Member State / Associate Country!!!	
E-AMPLE not to	



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### Validation result



The red 'Show Error' button indicates an error due to a missing or incorrect value in the form. The submission of the proposal **will be blocked** unless that specific field is corrected! Please complete the identified field.

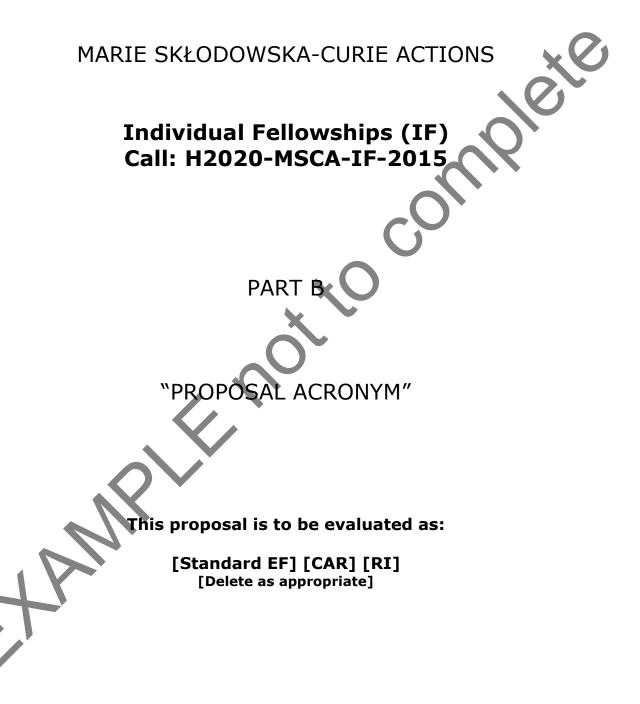
The yellow 'Show Warning' button indicates a warning due to a missing or incorrect value in the form. The submission of the proposal will not be blocked, the proposal will be submitted with the missing or incorrect value, which may cause problems at a later stage when processing the proposal. Please check and correct the identified field.

Section

#### Description

The form has not yet been validated, click "Validate Form" to do so!

### **START PAGE**



Part B - Page X of Y

#### TABLE OF CONTENTS

In drafting PART B of the proposal, applicants <u>must follow</u> the structure outlined below.

### LIST OF PARTICIPANTS START PAGE COUNT START PAGE COUNT STOP PAGE COUNT CV OF THE EXPERIENCED RESEARCHER CAPACITIES OF THE PARTICIPATING ORGANISATIONS COUNTAINS OF THE PARTICIPATING ORGANISATIONS

NB:

- Applicants must ensure that sections 1 3 do not exceed the limit of 10 pages.
- No reference to the outcome of previous evaluations of this or any similar proposal should be included in the text. Experts will be strictly instructed to disregard any such references.



#### List of Participants

Please provide a list of all participants (both beneficiary and, where applicable, partner organisations) indicating the legal entity, the department carrying out the work and the supervisor of the action.

If a secondment in Europe is planned but the partner organisation is not yet known, as a minimum the type of organisation foreseen (academic/non-academic) must be stated.

For non-academic beneficiaries, please provide additional data as indicated in the table below.

Participants	Legal Entity Short Name	Academic (tick)	non- academic (tick)	Countr Y	Dept./ Division / Laborator Y	Supervis or	Role of Partner Organisation <sup>1</sup>
<u>Beneficiary</u>							
- NAME						$\hat{\mathbf{O}}$	
Partner Organisation						5	
- NAME					xO		

#### Data for non-academic beneficiaries

Name	Location of research premises (city / country)	Type of R&D activities	No. ef full - time employees	No. of employees in R&D	Web site	Annual turnover (approx. in Euro)	Enterprise status (Yes/No)	SME status <sup>2</sup> (Yes/No)
		$\boldsymbol{X}$						

#### Note that:

- Any inter-relationship between different participating institutions or individuals (e.g. family ties, shared premises or facilities, joint ownership, financial interest, overlapping staff or directors, etc.) must be declared and justified in this part of the proposal;
  - The information in the table for non-academic beneficiaries **must be based on current data, not projections**;
- The data provided relating to the capacity of the participating institutions will be subject to verification during the Grant Agreement preparation phase.

**START PAGE COUNT** 

<sup>&</sup>lt;sup>1</sup> For example hosting secondments, for GF hosting the outgoing phase etc.

<sup>&</sup>lt;sup>2</sup> As defined in <u>Commission Recommendation 2003/361/EC.</u>

#### **1. Excellence**<sup>3</sup>

# **1.1** *Quality, innovative aspects and credibility of the research* (including inter/multidisciplinary aspects)

You should develop your proposal according to the following lines:

- Introduction, state-of-the-art, objectives and overview of the action
- <u>Research methodology and approach</u>: highlight the type of research and innovation activities proposed
- Originality and innovative aspects of the research programme: explain the contribution that the project is expected to make to advancements within the project field. Describe any novel concepts, approaches or methods that will be employed.

Explain how the high-quality, novel research is the most likely to open up the best career possibilities for the *Experienced Researcher* and new collaboration opportunities for the host organisation(s).

# **1.2** Clarity and quality of transfer of knowledge/training for the development of the researcher in light of the research objectives

Outline how a two way transfer of knowledge will occur between the researcher and the host institution, in view of their future development and past experience: (please see Section 5.2 of this Guide):

- Explain how the *Experienced Researcher* will gain new knowledge during the fellowship at the hosting organisation(s)
- Outline the previously acquired knowledge and skills that the researcher will transfer to the host organisation.

For Global Fellowships explain how the newly acquired skills and knowledge in the Third Country will be transferred back to the host institution in Europe during the incoming phase.

#### **1.3** Quality of the supervision and the hosting arrangements

Required sub-headings:

• Qualifications and experience of the supervisor(s)

Information regarding the supervisor(s) must include the level of experience on the research topic proposed and document their track record of work, including main international collaborations. Information provided should include participation in projects, publications, patents and any other relevant results.

 $<sup>^3</sup>$  Literature should be listed in footnotes, font size 8 or 9. All literature references <u>will count</u> towards the page limit.

#### • Hosting arrangements<sup>4</sup>

The text must show that the Experienced Researcher should be well integrated within the hosting organisation(s) in order that all parties gain the maximum knowledge and skills from the fellowship. The nature and the quality of the research group/environment as a whole should be outlined, together with the measures taken to integrate the researcher in the different areas of expertise, disciplines, and international networking opportunities that the host could offer.

For GF both phases should be described - for the outgoing phase, specify the practical arrangements in place to host a researcher coming from another country, and for the incoming phase specify the measures planned for the successful (re-)integration of the researcher.

Describe briefly how the host will contribute to the advancement of their career. In that context the following section of the European Charter for Researchers refers specifically to career development:

#### Career development

Employers and/or funders of researchers should draw up, preferably within the framework of their human resources management, a specific career development strategy for researchers at all stages of their career, regardless of their contractual situation, including researchers on fixed-term contracts. It should include the availability of mentors involved in providing support and guidance for the personal and professional development of researchers, thus motivating them and contributing to reducing any insecurity in their professional future. All researchers should be made familiar with such provisions and arrangements.

Therefore a Career Development Plan should not be included in the proposal, but it is part of implementing the project in line with the European Charter for Researchers.

# **1.4** Capacity of the researcher to reach and re-enforce a position of professional maturity in research

Applicants should demonstrate how their proposed research and personal experience can contribute to their professional development as an independent/mature researcher.

Please keep in mind that the fellowships will be awarded to the most talented researchers as shown by the proposed research and their track record (Curriculum Vitae, section 4), in relation to their level of experience.

<sup>&</sup>lt;sup>4</sup> The hosting arrangements refer to the integration of the Researcher to his new environment in the premises of the Host. It does not refer to the infrastructure of the Host as described in Criterion Implementation.

#### 2. Impact

# 2.1 Enhancing research- and innovation-related skills and working conditions to realise the potential of individuals and to provide new career perspectives

Explain the expected <u>impact of the planned research and training, and new</u> <u>competences acquired during the fellowship</u> on the capacity to increase career prospects for the Experienced Researcher after this fellowship finishes.

Demonstrate also to what extent competences acquired during the fellowship, including any secondments will increase the impact of the researcher's future activity on European society, including the science base and/or the economy.

# **2.2** Effectiveness of the proposed measures for **communication** and **results dissemination**

The new knowledge generated by the action should be used wherever possible to advance research, to foster innovation, and to promote the research profession to the public. Therefore develop following three points.

- <u>Communication and public engagement strategy of the action</u>
- Dissemination of the research results
- <u>Exploitation of results and intellectual property rights</u>

Concrete plans for the above must be included in the Gantt Chart (see point 3.1).

The following sections of the European Charter for Researchers refer specifically to public engagement and dissemination:

#### Public engagement

Researchers should ensure that their research activities are made known to society at large in such a way that they can be understood by non-specialists, thereby improving the public's understanding of science. Direct engagement with the public will help researchers to better understand public interest in priorities for science and technology and also the public's concerns.

#### Dissemination, exploitation of results

All researchers should ensure, in compliance with their contractual arrangements, that the results of their research are disseminated and exploited, e.g. communicated, transferred into other research settings or, if appropriate, commercialised. Senior researchers, in particular, are expected to take a lead in ensuring that research is fruitful and that results are either exploited commercially or made accessible to the public (or both) whenever the opportunity arises.

#### 3. Implementation

# **3.1 Overall coherence and effectiveness of the work plan,** including appropriateness of the allocation of tasks and resources

Describe the different work packages. The proposal should be designed in such a way to achieve the desired impact. A Gantt Chart should be included in the text listing the following:

- Work Packages titles (for EF there should be at least 1 WP);
- List of major deliverables;<sup>5</sup> <sup>6</sup>
- List of major milestones;<sup>7</sup>
- <u>Secondments if applicable.</u>

The schedule should be in terms of number of months elapsed from the start of the project.

# 3.2 Appropriateness of the management structure and procedures, including quality management and risk management

Develop your proposal according to the following lines:

- <u>Project organisation and management structure</u>, including the financial management strategy, as well as the progress monitoring mechanisms put in place;
- <u>Risks that might endanger reaching project objectives</u> and the contingency plans to be put in place should risk occur.

A deliverable is a distinct output of the action, meaningful in terms of the action's overall objectives and may be a report, a document, a technical diagram, a software, etc.

Deliverable numbers ordered according to 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.

<sup>7</sup> Milestones are control points in the action that help to chart progress. Milestones may correspond to the completion of a key deliverable, 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 action where, for example, the researcher must decide which of several technologies to adopt for further development.

PROPOSAL ACRONYM – Standard EF / CAR / RI (Delete as appropriate and include as header on each page)

**Example Gantt Chart** 

Reflecting work package, secondments, training events and dissemination / public engagement activities

Month	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24
Work package																								
Deliverable																			C					
Milestone																				-				
Secondment																	X	2						
Conference																X								
Workshop															)									
Seminar																								
Dissemination												V												
Public engagement																								
Other																								

CT ANN'

Delete rows and columns that do not apply.

#### 3.3 Appropriateness of the institutional environment (infrastructure)

- <u>Give a description of the main tasks</u> and commitments of the beneficiary and partners (if applicable).
- Describe the infrastructure, logistics, facilities offered in as far they are necessary for the good implementation of the action.

# **3.4 Competences, experience and complementarity** of the participating organisations and institutional commitment

The active contribution of the beneficiary to the research and training activities should be described. For GF also the role of partner organisations in Third Countries for the outgoing phase should appear. Additionally a letter of commitment shall also be provided in Section 7 (included within the PDF file of part B, but outside the page limit) for the partner organisations in Third Countries.

*NB:* Each participant is described in Section 5. This specific information should not be repeated here.

**STOP PAGE COUNT – MAX 10 PAGES** 

#### 4. CV of the Experienced Researcher

The CV is intrinsic to the evaluation of the whole proposal and is assessed throughout the 3 evaluation criteria.

This section should be limited to maximum 5 pages and should include **the standard academic and research record.** Any research career gaps and/or unconventional paths should be clearly explained so that this can be fairly assessed by the independent evaluators.

The *Experienced Researchers* must provide a list of achievements reflecting their track record, and this <u>may</u> include, <u>if applicable</u>:

- 1. **Publications** in **major**, **peer-reviewed conference proceedings and/or monographs** of their respective research fields, indicating also the number of citations (excluding self-citations) they have attracted.
- 2. Granted **patent(s)**.
- 3. **Research monographs, chapters** in collective volumes and any translations thereof.
- 4. **Invited presentations** to peer-reviewed, internationally established conferences and/or international advanced schools.
- 5. **Research expeditions** that the *Experienced Researcher* has led.
- 6. **Organisation of International conferences** in the field of the applicant (membership in the steering and/or programme committee).
- 7. Examples of participation in industrial innovation.
- 8. Prizes and Awards.
- 9. Funding received so far
- 10. Supervising, mentoring activities.

### 5. Capacity of the Participating Organisations

All organisations (whether beneficiary or partner organisation) must complete the appropriate table below, which will give input on the profile of the organisation as a whole. Complete one table of maximum <u>one page per institution, beneficiary or partner organisation</u> (min font size: 9). The experts will be instructed to disregard content above this limit.

Beneficiary X	0
General Description	×
Role and Commitment of key persons (supervisor)	(names, title, qualifications of the supervisor)
Key Research Facilities, Infrastructure and Equipment	Demonstrate that the team has sufficient facilities and infrastructure to host and/or offer a suitable environment for training and transfer of knowledge to recruited Experienced Researcher
Independent research premises?	Please explain the status of the beneficiary's research facilities – i.e. are they owned by the beneficiary or rented by it? Are its research premises wholly independent from other beneficiaries and/or partner organisations in the consortium?
Previous Involvement in Research and Training Programmes	Detail any relevant EU, national or international research and training projects in which the beneficiary has previously participated
Current involvement in Research and Training Programmes	Detail the EU and/or national research and training actions in which the partner is currently participating
Relevant Publications and/or research/innovation products	(Max 5) Produced by the organisation, not limited to the supervisor within the organisation.

Partner Organisation Y	
General description	
Key Persons and Expertise	
(supervisor)	
Key Research facilities,	
infrastructure and	
equipment	
Previous and Current	
Involvement in Research	
and Training Programmes	
Relevant Publications	(Max 3)
and/or research/innovation	
product	

### 6. Ethical Issues

Compliance with the relevant ethics provisions is essential from the beginning to the end of the project and is an integral part of research funded by the European Union within Horizon 2020.

Applicants submitting research proposals for funding within Marie Skłodowska-Curie actions in Horizon 2020 should demonstrate proactively to the REA that they are aware of and will comply with European and national legislation and fundamental ethical principles, including those reflected in the Charter of Fundamental Rights of the European Union<sup>8</sup> and the European Convention on Human Rights and its Supplementary Protocols.

Please be aware that it is the applicant's responsibility to identify any potential ethical issue, to handle the ethical aspects of the proposal and to detail how these aspects will be addressed.

#### The Ethics Review Procedure in Horizon 2020

All proposals above threshold and considered for funding will undergo an Ethics Review carried out by independent ethics experts. When submitting a proposal to Horizon 2020, all applicants are required to complete an **``Ethics Issues Table (EIT)**" in the Part A of the proposal. Applicants who flag ethical issues in the EIT have to also complete a more in depth **Ethics Self-Assessment in Part B**.

The ethics self-assessment will become part of the grant agreement and may thus lead to binding obligations that may later on be checked during ethics checks, reviews and audits.

For more details, please refer to the H2020 **"How to complete your Ethics Self- Assessment"** guide.

http://ec.europa.eu/research/participants/docs/h2020-funding-guide/crosscutting-issues/ethics\_en.htm

#### Ethics Self-Assessment (Part B)

The Ethics Self-Assessment must:

1) Describe how the proposal meets the EU and national legal and ethics requirements of the country/countries where the task raising ethical issues is to be carried out.

For more information on how to deal with Third Countries please see Article 34 of the annotated Model Grant Agreement<sup>9</sup>, as well as the following link:

http://ec.europa.eu/justice/data-protection/document/internationaltransfers/adequacy/index\_en.htm

Please list the documents provided with their expiry date.

<sup>&</sup>lt;sup>8</sup> The Charter of Fundamental Rights of the European Union: <u>http://www.europarl.europa.eu/charter/pdf/text\_en.pdf</u>

<sup>&</sup>lt;sup>9</sup> <u>http://ec.europa.eu/research/participants/data/ref/h2020/grants\_manual/amga/h2020-amga\_en.pdf</u>, page 235.

Ensure early compliance of the proposed research with EU and national legislation on ethics in research. Should your proposal be selected for funding, you will be required to provide as soon as possible the following documents (if applicable):

- an opinion from an Ethics Committee/Authority, required under national law;
- any other ethics-related documents mandatory under EU or national legislation;

If you have not already applied for/received the ethics approval/required ethics documents when submitting the proposal, please indicate in this section the approximate date when you will provide a missing approval/any other ethics documents, to the REA (scanned copy). Please state explicitly that you will not proceed with any research with ethical implications before the REA has received a scanned copy of all documents proving compliance with existing EU/national legislation on ethics.

If these documents are not issued in English, you are requested to submit also an English summary (containing in particular, if available, the conclusions of the Committee or Ethics Authority concerned).

If you plan to request these ethics documents specifically for your proposed project, your request must contain an explicit reference to the project's title.

# 2) Explain in detail how you intend to address the ethical issues flagged, in particular with regard to:

- the research **objectives** (e.g. study of vulnerable populations, cooperation with a Third Country, etc);
- the research **methodology** (e.g. clinical trials, involvement of children and related information and consent/assent procedures, data protection and privacy issues related to data collected, etc.);
- the potential **impact** of the research (e.g. dual use issues, environmental damage, malevolent use, etc.).





### **ENDPAGE**

### MARIE SKŁODOWSKA-CURIE ACTIONS



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