



# GUIDE FOR APPLICANTS

## *Information and Communication Technologies ICT*

Funding scheme:  
**COMBINATION OF COLLABORATIVE PROJECTS  
AND COORDINATION AND SUPPORT ACTIONS (CP-CSA)  
FOR PRE-COMMERCIAL PROCUREMENT (PCP)**

***FP7-ICT-2013-10***

*Further copies of this Guide, together with all information related to this Call for Proposals, can be downloaded via*

<http://ec.europa.eu/research/participants/portal/page/home>  
[http://cordis.europa.eu/fp7/ict/participating/home\\_en.html](http://cordis.europa.eu/fp7/ict/participating/home_en.html)

## About this Guide

This is version number 10 of the FP7 ICT Guide for Applicants for calls using single-stage submission procedures

Important information concerning the special conditions for Pre-Commercial Procurement proposals implemented by the combination of Collaborative projects and Coordination and Support actions (CP-CSA for PCP) appears throughout this Guide. Please read it with care!

*This Guide is based on the rules and conditions contained in the legal documents relating to FP7 (in particular the Seventh Framework Programme, Specific Programmes, Rules for Participation, and the Work programmes), all of which can be consulted via the CORDIS web-site. The Guide does not in itself have legal value, and thus does not supersede those documents.*

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## 1. Getting started

Funding decisions in the Seventh Framework Programme (FP7) are made on the basis of **proposals**. Proposals describe planned activities, information on who will carry them out, and how much they will cost. The Commission evaluates all eligible proposals in order to identify those whose quality is sufficiently high for possible funding. This **evaluation** is a peer-review carried out by independent experts.

The Commission then **negotiates** with some or all of those whose proposals have successfully passed the evaluation stage, depending on the budget available. If negotiations are successfully concluded, **grant agreements** providing for an EU financial contribution are established with the participants.

This **Guide for Applicants** contains the essential information to guide you through the mechanics of preparing and submitting a proposal. It is important that you have the correct Guide! Not only are there different Guides for different calls, there may also be different Guides for other funding schemes within the same call.

You must also refer to the current **ICT Workprogramme**. This provides a detailed description of the objectives and topics which are open for proposals, and will describe the wider context of research activities in this area. Work programmes are revised regularly, so make sure you refer to the latest version before preparing your proposal.

*Please check that this is the right guide for you by consulting the work programme and the **call fiche** (both documents posted on the CORDIS and on the Participant Portal websites), and the description of the funding scheme in the next section.*

This Guide and the work programme are essential reading. However, you may also wish to consult other reference and background documents, particular those relating to negotiation and the grant agreements, which will be made available on the Commission's CORDIS website:

[http://cordis.europa.eu/fp7/find-doc\\_en.html](http://cordis.europa.eu/fp7/find-doc_en.html) (see Annex 1 of this guide) and on the Participant Portal: <http://ec.europa.eu/research/participants/portal>.

All research activities supported by the Seventh Framework Programme should respect fundamental ethical principles.

## 2. About the funding scheme

### 2.1 General

A number of funding schemes are available to implement projects in FP7, but only certain ones may be available for the topics covered by this call. These are indicated in the call fiche.

This Guide covers a combination of Collaborative Projects and Coordination and Support Actions (CP-CSA) funding scheme for Pre-Commercial Procurement (PCP), and a description of this instrument is given later in this section. Please note that additional conditions may apply on a call-by-call basis. These will always be set out in the work programme (which includes the call fiche).

Note: Your proposal will be evaluated according to the funding scheme which you select. The Commission services will not re-examine or re-assign it on your behalf.

## **2.2 Combination of Collaborative Projects and Coordination and Support Actions (CP-CSA) for Pre-Commercial Procurement (PCP)**

### **Purpose**

The purpose of the action is to bring radical improvements to the quality and efficiency of public services with breakthrough solutions. This is done by encouraging public purchasers (such as national/regional/local governments, public transport operators, water or energy utilities,) around Europe to work together on new ICT based solutions that can respond to concrete public sector needs.

The public sector in the EU, as elsewhere in the world, is faced with important societal challenges such as climate change, energy and resource scarcity, an ageing population. Public sector transformations required to address those challenges are often so technologically demanding that no commercially stable solutions exist yet on the market, and new forward-looking public procurement strategies are needed that include the procurement of R&D.

Especially where interoperability and coherence of solutions across borders is required, cross-border cooperation can help better address issues of common European interest. To achieve the above targets, the action will support networking and cooperation between public bodies to define together the mid-to-long term solution requirements and to explore through PCP various alternative solution paths that respond to their needs.

This action aims to trigger a more forward-looking, concerted, public sector approach to societal challenges leading to:

- Cooperation between stakeholders across public sector departmental boundaries to develop common answers to societal challenges faced by the public sector across a number of EU Member or Associated States
- Reduced fragmentation of public sector demand by enabling public bodies to collectively implement PCP strategies in areas, which due to their nature are better addressed jointly, or which they would not have been able to tackle independently.
- Increased opportunities for wide market uptake and economies of scale for the supply side by forming critical mass on the public demand side, wide publication of results of cross border PCP activities and contribution to standardisation of jointly defined public sector PCP solution requirements specifications.

### **Size and resources**

The minimum number of participants is three mutually independent legal entities which are public purchasers preparing for or already experienced in the implementation of PCP. Each of these must be established in a different Member State or FP7 Associated Country (the countries concerned are listed in section 3 of this Guide). The entities must be independent of each other.

Eligible **public purchasers** include both contracting authorities in the meaning of the public procurement directive for public authorities (2004/18/EC) and utilities (2004/17/EC) at all levels (local, regional, national and supra-national) that plan to incorporate PCP in their public procurement programmes to improve the quality and efficiency of their public service offering. This includes for example public hospitals, public transport operators, relevant ministries (such as for health, welfare, transport, environment, justice, etc), water or energy utilities, communes, police or fire brigades, e-government administrations etc. Consortia shall demonstrate that they contain a critical mass of public purchasers - with clear financial commitments for jointly undertaking the cross-border PCP – that can trigger wide implementation of the public service innovation strategies and solutions that will be specified and/or developed during the PCP. The public purchasers are those that **would have to** purchase the new solutions that could be developed as a result of a pre-commercial procurement **in order to obtain** the desired quality and efficiency improvements in their public service offering. Please note that PCP only covers the procurement of R&D services, in

a way that is clearly separated from any potential subsequent purchases of large volumes of end-products (ref COM/2007/799). Participation to a PCP action therefore does not commit participating public purchasers to subsequent procurement of large volumes of end-products resulting from a PCP. Public purchasers undertaking a PCP can if they so desire, but are not obliged to, procure at market price R&D results resulting from a PCP (e.g. limited set of prototypes or test series products/services that were developed during a PCP).

**Other stakeholders**, that are important in the development of strategies for the innovation of public services through the implementation of PCP may participate in the action, if their participation is well justified and adds value to the action. This may be especially relevant for authorities or bodies with responsibility (e.g. related to certification, standardisation or regulation) in some area affected by the use of a particular novel technology that is addressed by the PCP. In some EU countries that have started creating policy programmes for encouraging innovation in public procurement, there are public authorities that co-organise/co-finance or provide financial incentives to public purchasers to undertake R&D procurements, such as PCP. Such public authorities are also eligible to participate in the CP-CSA for PCP consortia. These include for example those public authorities managing research and innovation programmes, those responsible for public procurement policy and/or structural funds programme management.

As CP-CSAs for pre-commercial procurement are targeted at public purchasers that will be actively involved in undertaking a joint PCP call for tender, **private bodies** that are **potential suppliers** of products or services in the area of solutions sought for by the public bodies through the PCP CP-CSAs are not eligible to participate as direct beneficiaries in the CP-CSA consortia. The reason for this being to exclude any conflict of interest, as one of the activities covered by the CP-CSA is the preparation of tender specifications and comparison/evaluation of competing solutions. Private bodies that are potential suppliers of solutions in the area sought for by the selected PCP CP-CSA consortia can only get involved after the CP-CSA is started. More specifically, the selected PCP CP-CSA consortia will launch an EU wide open call for tender to invite potential suppliers of solutions to send in offers for the research and development of the solutions sought for. During the preparation phase of the PCP call for tender, selected CP-CSA consortia are also requested to encourage an open dialogue<sup>1</sup> with potential tenderers and end-users to broach the views of the market about the R&D scope of the PCP.

**Private bodies** that are "**not**" **potential suppliers** of products or services in the area of solutions sought for by the public purchasers through the PCP CP-CSA, but whose participation is well justified, may participate in the action as direct beneficiaries (e.g. private hospitals, health insurance companies etc). With regards to the possibility for universities to participate in PCP projects, please note that the term industry also includes "for profit" universities. Because of the IPR sharing agreement to be used (see Appendix 6 of the WP), that assigns commercialisation responsibilities to the entities participating as solution developers in the pre-commercial procurement, the latter entities shall be "for-profit".

The size, scope and internal organisation of CP-CSAs for PCP can vary depending on the nature/complexity of R&D solution to be developed during the PCP, on the number of companies requested to participate in the PCP to have a good representation of possible competing solution paths, and on the composition of the consortium (number and level (local/regional/national) of public purchasers) needed to represent a sizeable critical mass on the demand side.

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<sup>1</sup> The open dialogue should be organised in a way not to preclude or distort competition. In respect of the Treaty principles, the open dialogue should be announced widely and well in advance and enable potential tenderers regardless of the geographic location to participate to the dialogue at least in English. All information given in answers to questions from participants in the dialogue should be documented and published.

## Duration

Pre-commercial procurement CP-CSAs are expected to last typically two to four years. However, there is no formal minimum or maximum duration.

## Activities

In their proposal for a CP-CSA for PCP, consortia shall have jointly identified a concrete challenge in the mid-to-long term innovation plans of the participating public purchasers that requires new R&D and that is proposed to be procured in cooperation through PCP. Consortia shall indicate in their proposal how the challenge, which is proposed as focus for the PCP, fits into their mid-to-long term innovation plans. Consortia shall describe in their proposal why solutions currently available on the market or under development are not meeting their needs, indicating that the level of ambition of the challenge proposed for the PCP is high enough to require new R&D to be undertaken. They shall put forward concrete targets for the desired functionality / performance improvement in the quality and efficiency of their public services that they would like to achieve by getting new solutions developed via the PCP, and SMART indicators that can be used to measure progress in achieving the targets.

Based on the action/implementation plan put forward in the proposal, the activities covered by a CP-CSA on PCP consequently include:

1. **CSA part: Networking and Coordination activities** for public bodies in Europe to cooperate in the innovation of their public services through a strategy that includes PCP.
2. **CP part: Joint research activities** related to validating the PCP strategy jointly defined by the public bodies participating in the action. This includes the exploration by the participating public bodies, through the execution of a joint pre-commercial procurement, of possible solutions for the targeted improvements in public sector services, and the testing of these solutions against a set of jointly defined performance criteria.

The two categories of activities are mandatory in the project due to the synergistic effects between the two components.

**1. Networking and Coordination activities** for public purchasers in Europe to cooperate in the innovation of their public services through a strategy that includes PCP, in particular activities related to preparation, management and coordination of the joint cross-border PCP undertaken under the CP part of the action, such as:

- Defining together the mid-to-long term solution requirements for the required public service innovation, and the resulting - not technology or solution prescriptive, but functional / performance based (see Appendix 6 of the WP) - specifications for a joint PCP call for tender
- Establishing and implementing good practice procedures for multinational PCP evaluation and monitoring (common evaluation criteria and implementation methods)
- Development of personnel exchange and/or joint training activities on PCP to help support a wider cooperation between public purchasers on introducing innovative solutions in public services across Europe (This may involve the allocation and training of additional resources in public bodies to develop a PCP implementation strategy.)
- Establishing specific cooperation agreements or (legal) arrangements between participants to prepare the ground for further trans-national PCP projects or programmes
- Dissemination of results and contribution to standardisation bodies (based on jointly defined requirements for the innovation of public services).
- Other networking and coordination activities essential to the preparation, management and coordination of a joint PCP call for tender, such as:
  - Building cooperation with other stakeholders essential for the implementation of a PCP strategy (e.g. between public purchasers, bodies responsible for R&D and innovation programmes, those responsible for developing public procurement policy and other stakeholders).

- Updating of the joint PCP action/implementation plan based on the outcomes of the PCP evaluations after each phase of the joint PCP. This involves the identification of common strategic issues essential for introducing the required innovations in public service provisioning, such as: points in time for cooperation with stakeholders, joint actions to be undertaken (e.g. regulatory agreements, joint standardisation activities) to ensure that potential obstacles are removed that hinder cross-border introduction of the innovative solutions under development.

**2. Joint research activities** related to validating the PCP strategy jointly defined by the public bodies participating in the action, through:

- Exploration, through a joint PCP, of possible alternative solution paths from a number of suppliers for the targeted improvements in public sector services, and
- Testing of these solutions against a set of jointly defined performance criteria, based on a well-defined public purchasing need that is jointly defined by the public purchasers undertaking the joint cross-border PCP.

Actions shall cover the full PCP life cycle of solution design, prototyping, and original development of a limited volume of products/services in the form of a test series.

The joint cross-border PCP involves the award of PCP contracts to a number of tenderers selected through a joint PCP call for tender organised during the action. Different constellations for joint procurement<sup>2</sup> are allowed, such as for example common procurement entity<sup>3</sup>, lead authority<sup>4</sup> and piggy-backing<sup>5</sup> constellations. A common mechanism, including a common set of selection/award criteria, for evaluating the offers submitted to the joint PCP call for tender shall be foreseen. Detailed rules for companies to participate in the financed projects shall be defined by the public purchasers. The call organisers shall organise the PCP while respecting the Treaty principles, the competition rules and the specific requirements in **Appendix 6** of the **work programme**. An important task of the selected consortia is to involve industry once the CP-CSA is started in an open, competitive and transparent way. As explained in more detail in Appendix 6 of the work programme, an EU wide published PCP call for tender shall be used to invite offers from all over Europe. Functional / performance based specifications shall be used in order to formulate the object of the PCP tender as a problem to be solved without prescribing a specific solution approach to be followed. Incoming offers will be evaluated not based on lowest price, but on best value for money (for more info see Appendix 6 of the work programme).

### **Financial Regime**

Reimbursement will be based on eligible costs (based on maximum rates of reimbursement specified in the grant agreement for different types of activities within the project). In some cases the reimbursement of indirect costs is based on a flat rate. The work programmes shall specify if other forms of reimbursement are to be used in the actions concerned. Participants in International Cooperation Partner countries (see Annex 1 of the work programme) may opt for a lump sum.

The EU contribution shall take the form of a CP-CSA grant that will combine

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<sup>2</sup> "Joint procurement" means combining the procurement actions of two or more contracting authorities into one procurement that is carried out jointly. The key defining characteristic for this action is that there should be only one tender published, and one entity awarding the resulting R&D service contracts for all PCP phases, on behalf of all participating contracting authorities.

<sup>3</sup> The "common procurement entity" constellation is an arrangement for joint procurement where all involved public purchasing authorities commonly establish or designate one external legal entity to conduct the joint procurement with a joint mandate and joint resources of all public purchasing authorities.

<sup>4</sup> The "lead authority" constellation is an arrangement for joint procurement where a group of public purchasing authorities collaborate through their existing purchasing departments in such a way that one public purchasing authority of the group is designated as lead authority to take responsibility for sourcing markets, tendering and arranging the R&D service contracts for all PCP phases, all in consultation with other purchasing authorities involved in the joint procurement.

<sup>5</sup> In the "piggy-backing" constellation one public purchasing authority executes the procurement and provides access to the results of the contract for a wider range of authorities, essentially by stating in the Contract Notice that other named public purchasing authorities may also wish to make use of the resulting contract a later date (normally during the timeframe of the original contract).



- (CSA part) reimbursement of 100% of eligible costs for the networking and coordination activities linked to the preparation, management and coordination of the joint PCP call for tender.<sup>6</sup> Reimbursement will be based on eligible costs (based on maximum rates of reimbursement specified in the grant agreement for different types of activities within the project). In some cases the reimbursement of indirect costs is based on a flat rate.
- (CP part) reimbursement of maximum 75% of the eligible costs<sup>7</sup> for the development of the new ICT solutions procured through the joint PCP call for tender (for financing the R&D to be performed by selected tenderers participating in the PCP).

CP-CSAs on PCP involve the organisation of a joint trans-national pre-commercial procurement by the beneficiaries of the CP-CSA grant. This involves the award of PCP contracts to tenderers selected through a joint PCP call for tender organised during the CP-CSA grant. Please note that, after award of the CP-CSA grant to the beneficiaries, the Commission will not be involved any more in the execution of the pre-commercial procurement undertaken jointly by the beneficiaries. Nevertheless, both for PCP CP-CSA proposals submitted in this call under objectives 5.1(d), 11.3 and 11.1<sup>8</sup>, the PCP procedure used by the beneficiaries shall be implemented according to the conditions in objective 11.1 and Appendix 6 of the ICT Work Programme 2013. If so provided in the call fiche, it is possible to claim subsistence costs and accommodation costs (related to travel as part of the implementation of a project) on the basis of flat rates. These rates, which do not cover travel costs, are in the form of a daily allowance for every country. The use of these rates is optional, but you may wish to use them when calculating your proposal budget. The rates themselves, and the detailed rules for their use, are given at this address: [http://cordis.europa.eu/fp7/find-doc\\_en.html](http://cordis.europa.eu/fp7/find-doc_en.html).

### **Specific Characteristics**

- The description of work (Annex I to the grant agreement) is normally fixed for the duration of the project. A sequence of updates of the description of work may be provided for in the grant agreement.
- The composition of the consortium is normally fixed for the duration of the project though enlargement of partnership, within the initial budget, is possible.

Proposers planning to submit a proposal for the CP-CSA instrument are **strongly recommended** to make use of the pre-proposal check service for this call which is described in Annex 1 of this Guide for applicants.

## **3. How to apply**

### ***3.1 Turning your idea into an effective proposal***

#### **The coordinator**

For a given proposal, the coordinator acts as the single point of contact between the participants and the Commission. The coordinator is generally responsible for the overall planning of the proposal and for building up the consortium that will do the work.

<sup>6</sup> The direct costs related to the CSA part of the action are funded at 100%.The reimbursement of indirect eligible costs related to the CSA part of the action may not exceed 7% of the direct eligible costs, excluding the direct eligible costs for subcontracting and the costs of reimbursement of resources made available by third parties which are not used on the premises of the participant.

<sup>7</sup> The maximum reimbursement rate is 75% for non-profit public bodies (e.g. for this action public purchasers that are subject to public procurement directive 2004/18/EC that are non-profit public bodies), secondary and higher education establishments or research organisations and SMEs which are unable to identify with certainty their real indirect costs for the project. The maximum reimbursement rate is 50% for other type of beneficiaries (e.g. for this action this includes private non-SME type public purchasers such as utilities that are subject to public procurement directive 2004/17/EC). It does not cover indirect costs (overheads).

<sup>8</sup> Within this call the objectives calling for CP-CSA proposals for PCP on specific topics in its area are : Objective ICT-2013.5.1(d) *Personalised health, active ageing and independent living* and Objective ICT-2013.11.3 on *High quality cloud computing environment for public sector needs*.

### Focusing your planned work

The work you set out in your proposal must correspond to one or more of the topics, and associated funding scheme(s), indicated in this call for proposals. Proposals that fail to do so will be regarded as ineligible.

*Multidisciplinary proposals addressing several topics may be submitted, provided that the 'centre of gravity' lies in a topic or topics open in the call in question.*

Refer to the Annex 2 of this Guide, and the work programme, to check the eligibility criteria and any other special conditions that apply. Refer also in those documents to the **evaluation criteria** against which your proposal will be assessed. Keep these in mind as you develop your proposal.

### Who can participate?

In principle, a legal entity may participate in a proposal no matter where it is established.

*A legal entity can be a so-called "natural person" (e.g. Mme Dupont) or a "legal person" (e.g. National Institute for Research).*

However, there are certain minimum conditions that have to be met relating to participation from the EU and Associated countries. These conditions vary between funding schemes (see section 2), and may also vary from call to call. See the call fiche for the conditions applicable to this call.

*The EU Member States are:*

*Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Netherlands, Poland, Portugal, Romania, Slovakia, Slovenia, Spain, Sweden, United Kingdom.*

*The Associated Countries are:*

*Albania, Bosnia and Herzegovina, Croatia, Faroe Islands, FYR Macedonia, Iceland, Israel, Liechtenstein, Moldova, Montenegro, Norway, Serbia, Switzerland, Turkey.*

*Other countries may become associated during the course of FP7. The latest news will be posted on the CORDIS web site.*

The following may receive EU funding in an FP7 project:

- Any legal entity established in a Member State or an Associated country (including the European Commission's Joint Research Centre), or created under Community law (e.g. a European Economic Interest Grouping),
- Any international European interest organisation (see Glossary).
- Any legal entity established in an FP7 International Cooperation Partner Country (ICPC). The list of ICPC can be found on the CORDIS web-site, and is given in Annex 1 of the work programme.
- Any other legal entity, under the conditions indicated below:
  - In the case of a participating international organisation, other than an international European interest organisation, or a legal entity established in a non-EU country other than an associated country or ICPC, a Community financial contribution may be granted provided that at least one of the following conditions is satisfied:
    - (a) Provision is made to that effect in the specific programmes or in the relevant work programme,
    - (b) It is essential for carrying out the indirect action,
    - (c) Such funding is provided for in a bilateral scientific and technological agreement or any other arrangement between the Community and the country in which the legal entity is established.

*Before the signature of a grant agreement, the Commission has to verify the existence and legal status of all participants. This verification is made only once for each organisation at the time of its first participation in FP7. The details of all validated organisations are stored in a **Unique Registration Facility (URF)**. These organisations are allocated a unique code, the so-called **Participant Identification Code (PIC)**. In any further participation in other proposals, the organisations already validated use the PIC for their identification with the Commission.*

*For the confirmation and maintenance of the data stored in the URF, the Commission asks each organisation to nominate one privileged contact person, the so-called Legal Entity Appointed Representative (LEAR). The LEAR is usually a person working in the central administration of the organisation and he/she must be appointed by the top management of the entity. The LEARs can view their organisations' legal and financial data online and ask for corrections and changes to the data of their legal entity via the Web interface of the Unique Registration Facility.*

### Cooperation with other countries

The Commission attaches great importance to international cooperation in research, and FP7 has been designed to ensure that such activities can be integrated across the programme. In addition to the opportunities mentioned above, which are generally applicable, calls may include:

- Topics of mutual interest defined in the work programmes where international cooperation is particularly encouraged.
- Specific international cooperation actions (SICA), also on topics of mutual interest. Here special minimum conditions apply.

Please check the work programme to see if these possibilities apply to this call.

More detailed practical advice on cooperation with third country participants in FP7 can be found here: [ftp://ftp.cordis.europa.eu/pub/fp7/docs/guideline-third-country-participants\\_en.pdf](ftp://ftp.cordis.europa.eu/pub/fp7/docs/guideline-third-country-participants_en.pdf)

### National Contact Points

A network of National Contact Points (NCPs) has been established to provide advice and support to organisations which are preparing proposals. You are highly recommended to get in touch with your NCP at an early stage. (see Annex 1 of this Guide).

Please note that the Commission will give the NCPs statistics and information on the outcome of the call (in particular, details of participants, but not proposal abstracts or funding details) and the outcome of the evaluation for each proposal. This information is supplied to support the NCPs in their service role, and is given under strict conditions of confidentiality.

### Other sources of help

Annex 1 of this Guide gives references to these further sources of help for this call. In particular:

- The Commission's general enquiry service on any aspect of FP7. Questions can be sent to a single e-mail address and will be directed to the most appropriate department for reply.
- The ICT Information Desk
- A dedicated help desk has been set up to deal with questions related to research ethics issues
- A dedicated help desk has been set up to deal with technical questions related to the Participant Portal Submission Service
- A further help desk providing assistance on intellectual property matters.
- Other services, including partner search facilities

### Proposal language

Proposals may be prepared in any official language of the European Union. If your proposal is not in English, a translation of the full proposal would be of assistance to the experts. An English translation of the abstract must be included in Part B of the proposal.

### Presenting your proposal

A proposal has two parts.

**Part A** will contain the administrative information about the proposal and the participants. The information requested includes a brief description of the work, contact details and characteristics of the participants, and information related to the funding requested (see Annex 3 of this Guide). This information will be encoded in a structured database for further computer processing to produce, for example, statistics and evaluation reports. This information will also support the experts and Commission staff during the evaluation process.

The information in Part A is entered through a set of on-line forms.

**Part B** is a "template", or list of headings, rather than an administrative form (see Annex 4 of this Guide). You should follow this structure when presenting the scientific and technical content of your proposal. The template is designed to highlight those aspects that will be assessed against the **evaluation criteria**. It covers, among other things, the nature of the proposed work, the participants and their roles in the proposed project, and the impacts that might be expected to arise from the proposed work.

Only black and white copies of Part B are used for evaluation and you are strongly recommended, therefore, not to use colour in your document. Do not insert hypertext links, only the text of your Part B will be read, not any documents linked to it.

Part B of the proposal is uploaded by the applicant into the **Participant Portal Submission Service**.

*A maximum length may be specified for the different sections of Part B, or for Part B as a whole (see Annex 4 of this Guide). You must keep your proposal within these limits. Information given on excess pages may be disregarded.*

*Even where no maximum page limits are given, it is in your interest to keep your text concise since over-long proposals are rarely viewed in a positive light by the evaluating experts.*

### Ethical principles

Please remember that research activities in FP7 should respect fundamental ethical principles, including those reflected in the Charter of Fundamental Rights of the European Union. Ethical principles include the need to ensure the freedom of research and the need to protect the physical and moral integrity of individuals and the welfare of animals. For this reason the European Commission carries out an ethical review of proposals when appropriate.

The following fields of research shall not be financed under this Framework Programme:

- research activity aiming at human cloning for reproductive purposes;
- research activity intended to modify the genetic heritage of human beings which could make such changes heritable<sup>9</sup>;

<sup>9</sup> Research relating to cancer treatment of the gonads can be financed.

- research activities intended 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.

As regards human embryonic stem cell research, the Commission will maintain the practice of the Sixth Framework Programme, which excludes from Community financial support research activities destroying human embryos, including for the procurement of stem cells. The exclusion of funding of this step of research will not prevent Community funding of subsequent steps involving human embryonic stem cells.

## 3.2 Proposal submission

### **About the Electronic Submission Services of the Commission**

Proposals must be submitted electronically, using the Commission's Electronic Submission Services which are to be found on the Participant Portal. Proposals arriving at the Commission/Agency by any other means are regarded as 'not submitted', and will not be evaluated<sup>10</sup>. All the data that the proposal coordinator uploads is securely stored on a server to which the proposal coordinator and the other participants in the proposal have access until after the call deadline.

The Electronic Submission Services can be accessed from the relevant call page on the Participant Portal.

As this is a web application, an Internet connection is required. An Internet browser and version 9 (or above) of the Adobe reader are needed. To check the requirements, click on <https://ec.europa.eu/research/participants/submission/manage/diagnostics>.

Full instructions are found in the "10 Minute guide to the Electronic Submission", available from the submission service website (click on "Starter Manual" to download the user guide).

### **Obtaining a mandatory Participant Identification Code (PIC):**

Before starting the process of submitting the proposal, **each participant in your proposal must be identified with a Participant Identification Code (PIC). Failure to do so will block the submission of your proposal!** The Participant Identification Code (PIC) is a unique 9 digit number that helps the Commission/Agency identify a participant organisation. It is used in all grant-related interactions between the organisation and the Commission/Agency. The use of PICs will lead to more efficient processing of your proposal.

If your organisation has already participated in a 7<sup>th</sup> Framework Programme proposal, it is likely that you already have a PIC number. You can check this on the Participant Portal: <http://ec.europa.eu/research/participants/portal/page/myorganisations> or in the search provided in the proposal submission system.

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<sup>10</sup> In exceptional cases, when a proposal co-ordinator has absolutely no means of accessing the Participant Portal Submission Service, and when it is impossible to arrange for another member of the consortium to do so, an applicant may request permission from the Commission to submit on paper. A request should be sent via the FP7 enquiry service (see annex 1), indicating in the subject line "Paper submission request". (You can telephone the enquiry service if web access is not possible: 00 800 6 7 8 9 10 11 from Europe; or 32 2 299 96 96 from anywhere in the world. A postal or e-mail address will then be given to you). Such a request, which must clearly explain the circumstances of the case, must be received by the Commission no later than one month before the call deadline. The Commission will reply within five working days of receipt. Only if a derogation is granted, a proposal on paper may be submitted by mail, courier or hand delivery. The delivery address will be given in the derogation letter.

*If your organisation already has a PIC, it is likely that it has also appointed a Legal Entity Authorised Representative (LEAR) (see section 3.1.). The names of LEARs are however not available online; you have to enquire within the administration of your own organisation.*

If a PIC is not yet available for an organisation, it can be obtained by registering the organisation in the Participant Portal under the 'Register' sub-tab of the 'My Organisations' tab (<http://ec.europa.eu/research/participants/portal/page/myorganisations>). After filling in the data for the entity, a PIC number is given, which can then be used in the Electronic Submission Services. [You are encouraged to proceed well before the call deadline to avoid potential last minute troubles.](#)

All participants already possessing a PIC should use it to identify themselves in the Electronic Submission Service. After entering the PIC, sections of their A forms are filled in automatically.

If, after entering your PIC, the data which appears for your organisation is incorrect, you should contact the LEAR of your organisation to correct it through the Participant Portal. You can also change yourself the data prefilled from the PIC in your A forms, but these changes will remain local to the Electronic Submission Services for this proposal only. The original data, which is stored in the Commission's database, will always re-appear whenever the PIC is used, until the new data is validated.

## **Get started**

As a first step, the coordinator starts creating the proposal by accessing the system from the call page. Access to the Electronic Submission Service is granted after logging in to the Participant Portal from the relevant call's page.

### **Proposal coordinator's actions:**

#### **Step one: getting a European Commission Authentication Service (ECAS) user ID..**

Getting a personal user ID with the **European Commission Authentication Service (ECAS)** is mandatory in order to login to the Participant Portal and to be able to use the different functions of the Portal, including the proposal submission. This will allow editing the proposal data in the electronic proposal submission system, completing the information requested or – for coordinators - submitting the proposal. The system will request a login by every partner. The same user ID will be used for all later interactions with the Commission/Agency in the field of Research. [Further details on the ID are available under: https://webgate.ec.europa.eu/cas/eim/external/help.cgi.](https://webgate.ec.europa.eu/cas/eim/external/help.cgi)

#### **Step two: choosing a funding scheme**

For each call, a list of available funding schemes or objectives as activity codes will be presented by the Electronic Submission Services. The proposal coordinator must choose the appropriate one for the proposal. Refer to the call fiche and work programme for the various conditions applicable to each funding scheme.

#### **Step three: creating a draft proposal**

Once the coordinating organisation is identified with its PIC number, the coordinator fills in the pre-registration data for the proposal: acronym, short summary, activity code, and at the next step, the list of participants. These details can be used by the Commission/Agency services in order to plan the evaluation. In general, the following details are requested:

- *The proposal acronym.* This is the name of the proposal and it will be used throughout the lifetime of the project, if funded. No more than 20 characters are allowed (standard alphabet and numbers only; no symbols or special characters, except underscore, space, hyphen or dot).
- *The proposal short summary,* which describes briefly the purpose of the proposal with a maximum of 2,000 characters. Entering at least keywords will help the services in preparing the evaluations (e.g. choosing the experts for the evaluations). Coordinators may choose to enter 'xxx' at this stage should they prefer not disclosing any data.
- *Activity code,* the objective addressed by the proposal.

#### **Step four: adding other participants to the proposal**

At this step the proposal coordinator sets up the consortium. The proposal coordinator can:

- *Add other participants to the proposal.*
  - The coordinator adds the partners using the nine-digit identifier, the PIC number. A search function is provided to help the coordinator finding the PIC number of the partners.
  - Once the coordinator has added the entities to the consortium, the coordinator has to insert the contact persons' details for each participant. The main identifier is the e-mail address of a person.
  - Once the coordinator saves this page, an automatic invitation is sent to all contacts' e-mail address. The invited persons can access the proposal after logging in to the Participant Portal - with the ECAS account linked to the given e-mail address - under the 'My Proposals' tab.
- *Delete a participant.*
- *Reorder the participants.* The order of the participants in the administrative forms will be adjusted.

The proposal coordinator however cannot be deleted, and is always the first participant.

#### **Step five: forms, files and submission**

This step is the core of the process, as, from this step, the proposal coordinator can:

- Fill in the administrative forms, part A of the proposal (see Annex 3 of this Guide)
- Forms are completed using a PDF reader (e.g. Adobe Reader, see above "What is needed" for minimum requirements). The proposal coordinator can complete all the forms, including the budget table and the administrative details of the coordinator and of all participants. Proposal partners can only complete their own administrative details (form A2).
- Download the template of the part B of the proposal and other information files (see Annex 4 of this Guide)
- Upload the file that will be the part B of the proposal.
- Submit the proposal package.

**Only the coordinator can upload the part B of the proposal and submit the proposal. Therefore, only the coordinator should be logged into the Electronic Submission Service when the submission attempt is made.**

For the proposal Part B you must use exclusively PDF ("portable document format", compatible with Adobe version 3 or higher, with embedded fonts). Other file formats will not be accepted by the system. Irrespective of any page limits specified in annex 4 to this Guide, there is an overall limit of 10 Mbytes to the size of proposal file Part B. It is advised to limit the size of the proposal to 2 Mbytes.

There are also restrictions to the name given to the Part B file: use alphanumeric characters; special characters and spaces must be avoided.

*You are advised to clean your document before converting it to PDF (e.g. accept all tracked changes, delete notes).*

*Check that your conversion software has successfully converted all the pages of your original document (e.g. there is no problem with page limits).*

*Check that your conversion software has not cut down landscape format pages to fit them into portrait format. Check that captions and labels have not been lost from your diagrams*

*Please note that the Commission prints out proposals in black and white on plain A4 paper. The printable zone on the print engine is bounded by 1.5 cm right, left, top bottom. No scaling is applied to make the page "fit" the window. Printing is done at 300 dots per inch.*

Completing the Part A forms in the Electronic Submission Services and uploading a Part B does not yet mean that the proposal is submitted. Once there is a consolidated version of the proposal, the "SUBMIT" button must be pressed. **Only the coordinator is authorised to submit the proposal.**

At this point the service performs a limited automatic validation of the proposal. A list of discovered problems, such as missing data, is given on the last page of the proposal submission forms. In some cases users are allowed to submit incomplete administrative information **but for significant omissions, proposal submission will be blocked until the problems are corrected.** Therefore you are strongly advised, when preparing your proposal, to regularly click on 'validate' at the bottom of any page of the Part A to obtain updated validation messages and to review them on the last page of the proposal submission forms.

When errors or omissions are corrected, the coordinator must then repeat the above step to finally achieve the proposal submission.

If the submission sequence described above is not followed,  
the Commission/Agency considers that no proposal has been submitted.

When the proposal has been successfully submitted, the service will proceed to Step 6 where the coordinator sees a message that indicates that the proposal has been received.  
(This automatic message is not the official acknowledgement of receipt - see Section 5).

#### **Step six: proposal status page**

Reaching this step means that the proposal is submitted (i.e. sent to the Commission/Agency services for evaluation). It does not mean that the proposal is valid, complete, eligible in all respects or that it will be funded.

In Step six you can:

- *Download the proposal.* It is advised to download the proposal once submitted to check that it has been correctly sent. The downloaded proposal will be digitally signed and time stamped.



- *Re-edit the proposal*, going back to step 5. After re-editing the proposal (any data in the forms or a modified attachment), modifications have to be resubmitted. The coordinator may continue to modify the proposal and submit revised versions overwriting the previous one right up until the deadline.
- *Withdraw the proposal*. If the proposal is withdrawn, it will not be considered for evaluation. A reason for the withdrawal will be requested by the service.  
(Note: Your proposal draft is not deleted from the server and this withdrawal action can be reversed, but only before the deadline, by simply submitting it again).

### **Use of the system by the other participants**

In order to access the proposals, all contacts need to have an ECAS ID that is necessary for the login of the Participant Portal.

Those contacts, who have been invited by the coordinator, can access the proposal via the 'My Proposals' tab after login.

In the proposal submission system, participants can

- complete their own entity's administrative details and budget forms (A2 and A3.1);
- download the document template for writing Part B of the proposal, in order to assist the coordinator in preparing it, however, only the coordinator can upload the finished version;
- view the whole proposal.

### **About the deadline**

Proposals must be submitted on or before the deadline specified in the call fiche. It is your responsibility to ensure the timely submission of your proposal.

The Electronic Submission Services of the Commission will be closed for this call at the call deadline. Please note that the deadline established in the call refers to the sharp time, no extra seconds allowed (for example, call deadline at 17.00 means at 17.00.00). After this moment, the proposal can no longer be modified. It is however visible in a read-only version.

***Do not wait until the last moment before submitting your proposal!  
Call deadlines are absolutely firm and are strictly enforced.***

Please note that successive drafts of the proposal can be submitted with the Electronic Submission Services. Each successive submission overwrites the previous version. It is a good idea to **submit a draft well before the deadline**.

*Do not wait until the last moment to attempt the submission of the proposal. Internet access issues and proposal verification issues must be detected well before the submission deadline if help is to be requested from the service desk. Such issues are never accepted as extenuating circumstances for failure to submit in time.*

*Submission is deemed to occur at the moment when the proposal coordinator completes the submission sequence described above. It is not the point at which the upload of the part B is started. If you wait until too near to the close of the call to start uploading your proposal, there is a serious risk that you will not be able to submit in time.*

*If you have submitted your proposal in error to another call which closes after this call, the Commission/agency will not be aware of it until it is discovered among the downloaded proposals for the later call. It will therefore be classified as ineligible because of late arrival.*

*The submission of a proposal requires some knowledge of the Electronic Submission Services, a detailed knowledge of the contents of the proposal and the authority to make last-minute decisions on behalf of the consortium if problems arise. **You are advised not to delegate the job of submitting your proposal!***

In the unlikely event of a failure of the Electronic Submission Services due to breakdown of the Commission server during the last 24 hours of this call, the deadline will be extended by a further 24 hours. This will be notified by e-mail to all proposal coordinators who had registered for this call by the time of the original deadline, and also by a notice on the Call pages on the Participant Portal. Such a failure is a rare and exceptional event; therefore do not assume that there will be an extension to this call. If you have difficulty in submitting your proposal, you should not assume that it is because of a problem with the Commission servers, as this is rarely the case. Contact the Electronic Submission Services help desk if in doubt (see the address given in annex 1 of this Guide).

Please note that the Commission/Agency will not extend deadlines for system failures that are not its own responsibility. In all circumstances, you should aim to submit your proposal well before the deadline to have time to solve any problems.

*A small number of calls operate a **continuous submission procedure**. These calls are open for an extended period, during which proposals will be evaluated in batches after fixed cut-off dates. The call fiche will show whether intermediate cut-off dates apply to his call.*

### **Correcting or revising your proposal**

Errors discovered in proposals submitted can be rectified by simply submitting a corrected version before the submission deadline; the new proposal package (part A and B) will overwrite the old one.

Once the deadline has passed, however, the Commission/Agency can accept no further additions, corrections or re-submissions.

**The last version of your proposal submitted before the deadline is the one which will be taken into consideration; no later version can be substituted and no earlier version can be recovered.**

### **Ancillary material**

Only a single PDF file comprising the complete Part B can be uploaded. Unless specified in the call, any hyperlinks to other documents, embedded material, and any other documents (company brochures, supporting documentation, reports, audio, video, multimedia etc.) sent electronically or by post will be disregarded.

### **Withdrawing a proposal**

You may withdraw a proposal before the call deadline by simply submitting a revised version with an empty Part B section, and with the following text in the abstract field of form A1:

*"The applicants wish to withdraw this proposal. It should not be evaluated by the Commission".*

You may also withdraw a proposal, either before or after the call deadline, by accessing the 'My Proposals' tab when you log in to the participant portal. With the action 'view submitted' the coordinator will move to Step 6, where the proposal can be withdrawn.

A withdrawn proposal will not be subsequently considered by the Commission.

### **Registration of legal entities in the Commission's Early Warning System (EWS) and Central Exclusion Database (CED).**

To protect the EU's financial interests, the Commission/Agency uses an internal information tool, the Early Warning System (EWS) to flag identified risks related to beneficiaries of centrally managed contracts and grants. Through systematic registration of financial and other risks the EWS enables the Commission services to take the necessary precautionary measures to ensure sound financial management<sup>11</sup>.

EWS registrations are not publicly disclosed. However, registrations will be transferred to the Central Exclusion Database (CED) if they relate to entities that have been excluded from EU funding because they are insolvent or have been convicted of serious professional misconduct or a criminal offence detrimental to EU financial interests. The data in CED are available to **all public authorities implementing EU funds**, i.e. European institutions, national agencies or authorities in Member States, and, subject to conditions for personal data protection, to third countries and international organisations.

The work programme informs you that the details of your organisation (or those of a person who has powers of representation, decision-making or control over it) may be registered in the EWS and the CED and be shared with public authorities as described in the relevant legal texts<sup>12</sup>.

More information on the EWS and CED can be found here:

[http://ec.europa.eu/budget/explained/management/protecting/protect\\_en.cfm](http://ec.europa.eu/budget/explained/management/protecting/protect_en.cfm)

### **Data protection**

Proposals are archived under secure conditions at all times. The data contained in the proposal are treated in accordance with Regulation 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<sup>13</sup>. After completion of the evaluation and at any subsequent negotiation, all copies are destroyed other than those required for archiving and/or auditing purposes.

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<sup>11</sup> The EWS covers situations such as significantly overdue recovery orders, judicial proceedings pending for serious administrative errors/fraud, findings of serious administrative errors/fraud, legal situations which exclude the beneficiary from funding.

<sup>12</sup> The basis of registrations in EWS and CED is laid out in: - the Commission Decision of 16.12.2008 on the Early Warning System (EWS) for the use of authorising officers of the Commission and the executive agencies (OJ, L 344, 20.12.2008, p. 125), and- the Commission Regulation of 17.12.2008 on the Central Exclusion Database – CED (OJ L 344, 20.12.2008, p. 12).

<sup>13</sup> (OJ L 8, 12.1.2001).

## 4. Check list

### 4.1 Preparing your proposal

- **Does your planned work fit with the call for proposals?** Check that your proposed work does indeed address the topics open in this call. (See the current version of the work programme).
- **Are you applying for the right call and funding scheme?** Check that you have applied for the right call and one of the funding schemes open for your chosen topic (see the work programme)<sup>14</sup>.
- **Is your proposal eligible?** The eligibility criteria are given in the work programme. See also Annex 2 of this Guide. In particular, make sure that you satisfy the minimum requirements for the makeup of your consortium. Have any additional eligibility criteria been set for this call? Check that you comply with any budgetary limits that may have been fixed on the requested EU contribution. Any proposal not meeting the eligibility requirements will be considered ineligible and will not be evaluated.
- **Is your proposal complete?** Proposals must comprise a Part A, containing the administrative information including participant and project cost details on standard forms; and a Part B containing the scientific and technical description of your proposal as described in this Guide. A proposal that does not contain both parts will be considered ineligible and will not be evaluated.
- **Does your proposal follow the required structure?** Proposals should be precise and concise, and must follow exactly the proposal structure described in this document (see Annex 4 of this Guide), which is designed to correspond to the evaluation criteria which will be applied. Omitting requested information will almost certainly lead to lower scores and possible rejection.
- **Does your proposed work raise ethical issues?** Clearly indicate any potential ethical, safety or regulatory aspects of the proposed research and the way these will be dealt with prior and during the implementation of the proposed project. A preliminary ethics control will take place during the evaluation and, if needed, an ethics screening and/or review will take place for those proposals raising ethics issues. Proposals may be rejected on ethical grounds if such issues are not dealt with satisfactorily.
- **Have you maximised your chances?** There will be strong competition. Therefore, edit your proposal tightly, strengthen or eliminate weak points. Put yourself in the place of an expert evaluator; refer to the evaluation criteria given in Annex 2 of this Guide. Arrange for your draft to be evaluated by experienced colleagues; use their advice to improve it before submission.
- **Do you need further advice and support?** You are strongly advised to inform your National Contact Point of your intention to submit a proposal (see address in Annex 1 of this Guide). Remember also the Enquiry service listed in Annex 1 of this Guide.

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<sup>14</sup> If you have in error registered for the wrong call or funding scheme, discard that registration (usernames and passwords) and register again before the call deadline. If, after the close of the call, you discover that you have submitted your proposal to the wrong call, notify the Participant Portal Submission Service Helpdesk.

## **4.2. Final checks before submission**

- Do you have the agreement of all the members of the consortium to submit this proposal on their behalf?
- Is your Part B in portable document format (PDF), including no material in other formats?
- Is your Part B filename made up only of the letters A to Z and numbers 0 to 9 without special characters or spaces?
- Have you printed out your Part B, to check that it really is the file you intend to submit, and that it is complete, printable and readable? After the call deadline it will not be possible to replace your Part B file
- Have you respected the font size (11 point) and the page limitations for the different sections?
- Is your Part B file within the size limit of 10 Mbytes?
- Have you virus-checked your computer? The Participant Portal Submission Service will automatically block the submission of any file containing a virus.

## **4.3. The deadline: very important!**

- Have you made yourself familiar with the Participant Portal Submission Service in good time?
- Have you allowed time to submit a draft version of your proposal well in advance of the deadline (at least several days before), and then to continue to improve it with regular resubmissions?
- Have you completed the Participant Portal submission process for your final version?

## **4.4. Following submission**

- Proposals submitted to the Participant Portal can still be reviewed by the applicant.
- It is highly recommended that after uploading and submitting your final version, you then review what you have uploaded
- Do this while there is still time to submit a corrected version if necessary

## **5. What happens next**

Shortly after the call deadline (or batch date in the case of continuously open calls), the Commission will send an **Acknowledgement of receipt** to the e-mail address of the proposal coordinator given in the submitted proposal. This is assumed to be the individual named as “person in charge” on the A2 form of participant no. 1. Please note that the brief electronic message given by the Participant Portal Submission Service after each submission is not the official Acknowledgement of receipt.

The sending of an acknowledgement of receipt does not imply that a proposal has been accepted as eligible for evaluation.

*If you have not received an Acknowledgement of receipt within 12 working days after the call deadline (or cut-off date, in the case of a continuously open call), you should contact the FP7 Enquiry Service (see Annex 1 of this Guide). However, first please check that you are the person named in the proposal as contact person for partner no. 1, check the email address which you gave for*

*yourself, and check the junk mail box of your email system for a few days following the close of call for any mail originating from [FP7Aor@ess-fp7.org](mailto:FP7Aor@ess-fp7.org).*

The Commission will check that your proposal meets the **eligibility criteria** that apply to this call and funding scheme (see the work programme and Annex 2 of this Guide).

All eligible proposals will be evaluated by independent experts. The evaluation criteria and procedure are described in Annex 2 of this Guide.

Soon after the completion of the evaluation, the results will be finalised and all coordinators will receive a letter containing initial information on the results of the evaluation. However, even if the experts viewed your proposal favourably, the Commission cannot at this stage indicate if there is a possibility of EU funding.

*If you have not received your ESR by the date referred to in Annex I of this Guide, please contact the Commission via the FP7 enquiry service.*

The letter will also give the relevant contact details and the steps to follow if you consider that there has been a shortcoming in the conduct of the evaluation process ("redress procedure").

The Commission also informs the relevant **programme committee**, consisting of delegates representing the governments of the Member states and Associated countries.

Based on the results of the evaluation by experts, the Commission draws up the final list of proposals for possible funding, taking account of the available budget.

Official letters are then sent to the applicants. If all has gone well, this letter will mark the beginning of a **negotiation** phase. Due to budget constraints, it is also possible that your proposal will be placed on a reserve list. In this case, negotiations will only begin if funds become available. In other cases, the letter will explain the reasons why the proposal cannot be funded on this occasion.

Negotiations between the applicants and the Commission aim to conclude a grant agreement which provides for EU funding of the proposed work. They cover both the scientific/technological, and the administrative and financial aspects of the project. The officials conducting these negotiations on behalf of the Commission will be working within a predetermined budget envelope. They will refer to any recommendations which the experts may have made concerning modifications to the work presented in the proposal, as well as any recommendations arising from an ethical review of the proposal if one was carried out. Where relevant, security aspects shall also be considered.

The negotiations will also deal with gender equality actions, and, if applicable to the project, with gender aspects in the conduct of the planned work, as well as the relevant principles contained in the European Charter for researchers and the Code of Conduct for their recruitment.

A description of the negotiation process will be provided in the "**FP7 Guidelines for negotiation notes**" (to be made available on CORDIS). Members of the proposal consortium may be invited to Brussels or Luxembourg to facilitate the negotiation.

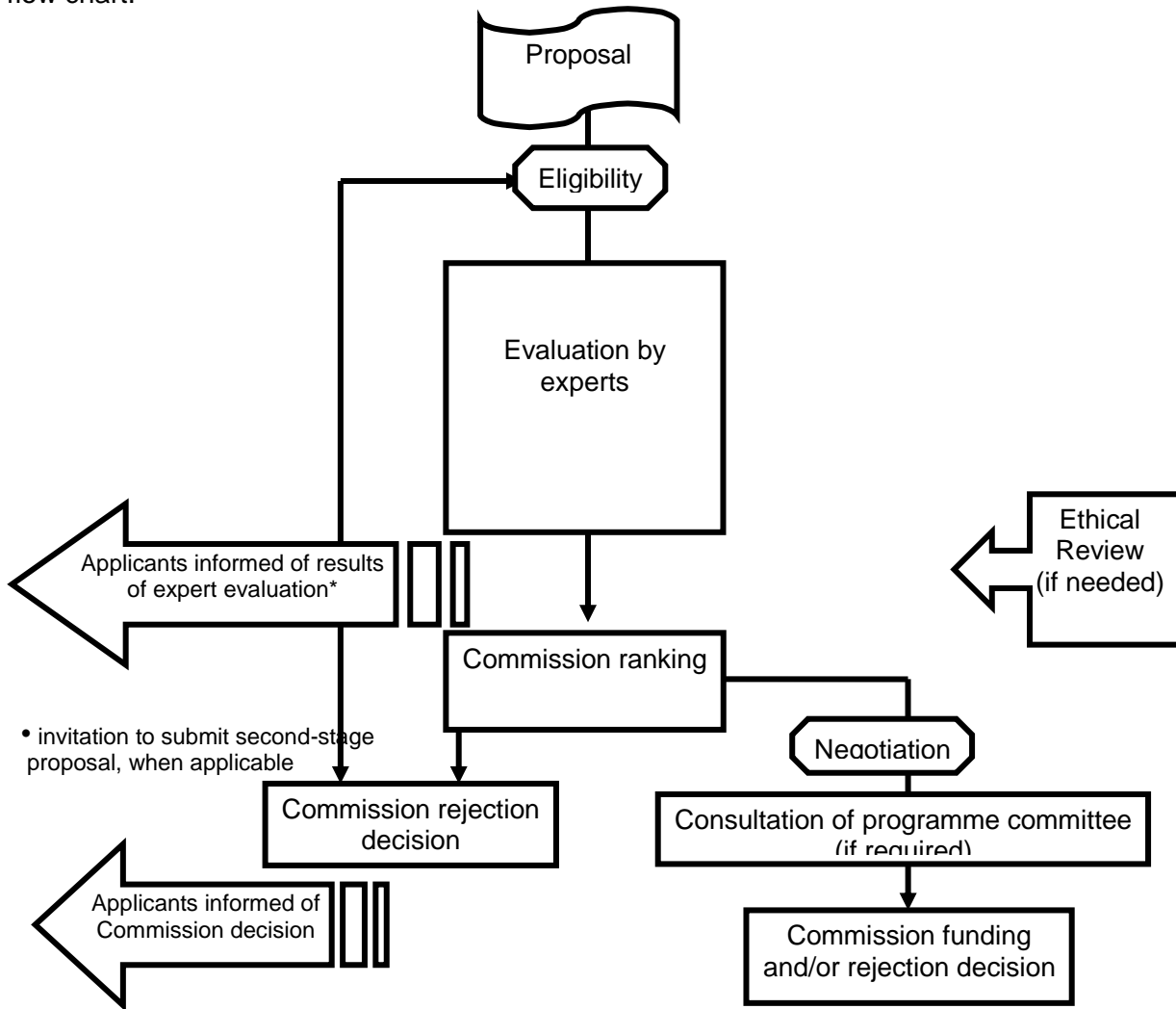
For participants not yet having a Participant Identification Code (PIC), i.e. not yet being registered and validated in the Commission's Unique Registration Facility (URF) their existence as legal entities and their legal status will have to be validated before a grant agreement can be signed.

Applicants are reminded that the Commission's Research DGs have adopted a new and reinforced audit strategy aimed at detecting and correcting errors in cost claims submitted in projects on the basis of professional auditing standards. As a result the number of audits and participants audited

will increase significantly and the Commission's services will assure appropriate mutual exchange of information within its relevant internal departments in order to fully coordinate any corrective actions to be taken in a consistent way. More information can be found here: [http://cordis.europa.eu/audit-certification/home\\_en.html](http://cordis.europa.eu/audit-certification/home_en.html)

Summary of the evaluation and selection process

The sequence of steps in the evaluation and selection procedure is summarised in the following flow chart:



Risk Sharing Finance Facility (RSFF)

This innovative debt-based facility, designed by the European Commission and the European Investment Bank, creates an additional capacity of up to €10 billion for financing higher risk research, technological development, demonstration and innovation activities.

The EIB will implement RSFF in close collaboration with all major EU national and regional banks within Member states and Associated countries to FP7, which are providing support to the development of European companies.

Financing through the RSFF can be sought either in addition to, or instead of FP7 grants.

For additional information on RSFF see:

<http://www.eib.org/products/loans/special/rsff/index>

[http://ec.europa.eu/invest-in-research/funding/funding02\\_en.htm](http://ec.europa.eu/invest-in-research/funding/funding02_en.htm)



## Glossary

The following explanations are provided for clarity and easy-reference. They have no legal authority, and do not replace any official definitions set out in the Council decisions.

### A

Acknowledgement of receipt :

Applicants are informed by email shortly after the deadline that a proposal has been successfully submitted (but not that it is necessarily eligible). Contact the FP7 Enquiry service urgently if you do not receive such an acknowledgement within a few days of the close of call (or batch, for continuous submission calls).

Applicant

The term used generally in this guide for a person or entity applying to a call for proposals. The term 'participant' is used in the more limited sense of a member of a proposal or project consortium (see below).

Associated countries

Non-EU countries which are party to an international agreement with the Community, under the terms or on the basis of which it makes a financial contribution to all or part of the Seventh Framework Programme. In the context of proposal consortia, organisations from these countries are treated on the same footing as those in the EU. The list of associated countries is given in the body of this guide.

### C

Call fiche

The part of the work programme giving the basic data for a call for proposals (e.g. topics covered, budget, deadline etc). It is posted as a separate document on the Participant Portal web pages devoted to a particular call.

Call for proposals (or "call")

An announcement is published, usually in the Official Journal, inviting proposals for research activities in a certain theme. Full information on the call can be found on the Participant Portal web-sites.

Consensus meeting

The stage, in the proposal evaluation process, when experts come together to establish a common view on a particular proposal.

Consortium

Most funding schemes require proposals from a number of participants (usually at least three) who agree to work together in a consortium.

Continuous submission

Some calls are open for an extended period, during which proposals may be submitted at any moment. In these cases, proposals are evaluated in batches after fixed cut-off dates.

Coordinator

The coordinator leads and represents the applicants. He or she acts as the point of contact with the Commission.

#### CORDIS service

A web service providing access to all the documentation related to FP7. (See also Participant Portal).

#### Cut-off date

An intermediate date in the context of a call operating a continuous submission procedure. Proposals are evaluated in batches after each cut-off date.

### **D**

#### Deadline

For a particular call, the moment after which proposals cannot be submitted to the Commission, and when the Participant Portal Submission Service closes for that call. Deadlines are strictly enforced.

#### Deliverable

A deliverable represents a verifiable output of the project. Normally, each workpackage will produce one or more deliverables during its lifetime. Deliverables are often written reports but can also take another form, for example the completion of a prototype etc.

#### Direct costs

Direct costs are all eligible costs which can be attributed directly to the project and are identified by the participant as such, in accordance with its accounting principles and its usual internal rules.

### **E**

#### Early Warning System (EWS)

An internal information tool of the Commission to flag identified financial risks related to beneficiaries.

#### Eligibility committee

An internal committee which examines in detail cases of proposals whose eligibility for inclusion in an evaluation is in question

#### Eligibility criteria

The minimum conditions which a proposal must fulfil, if it is to be retained for evaluation. The eligibility criteria are generally the same for all proposals throughout FP7, and relate to submission before the deadline, minimum participation, completeness and scope. However, additional eligibility criteria may apply to certain calls, and applicants should check the work programme, and annex 2 to this Guide.

#### Ethical issues table

Research activities supported by the Framework Programme should respect fundamental ethical principles. The main issues which might arise in a project are summarised in tabular form in a checklist included in the proposal

#### Evaluation criteria

The criteria, against which eligible proposals are assessed by independent experts. The evaluation criteria are generally the same for all proposals throughout FP7, and relate to S/T quality, impact and implementation. Relevance is also considered. However, additional evaluation criteria may apply to certain calls, and applicants should check the work programme, and annex 2 to this Guide.

#### Evaluation Summary Report (ESR)

The assessment of a particular proposal following the evaluation by independent experts is provided in an Evaluation Summary Report. It normally contains both comments and scores for each evaluation criterion.

## **F**

FP7 enquiry service

A general information service on all aspects of FP7. Contact details are given in annex 1 to this Guide.

Funding scheme

The mechanisms for the Community funding of research projects. The funding schemes have different objectives, and are implemented through grant agreements.

## **G**

Grant Agreement (GA)

The legal instrument that provides for Commission funding of projects.

## **H**

Hearing

Applicants whose proposals have been evaluated are sometimes invited to provide explanations and clarifications to any specific questions raised by the experts. These questions are transmitted to the applicants in advance.

## **I**

Indirect costs

Indirect costs, (sometimes called overheads), are all those eligible costs which cannot be identified by the participant as being directly attributed to the project, but which can be identified and justified by its accounting system as being incurred in direct relationship with the eligible direct costs attributed to the project.

Individual evaluation

The stage in the evaluation process, when experts assess the merits of a particular proposal before discussion with their peers.

Information Days

Open events organised by the Commission to explain the characteristics of specific calls, and often as well, a chance for potential applicants to meet and discuss proposal ideas and collaborations.

Initial information letter

The letter sent by the Commission to applicants shortly after the evaluation by experts, which includes the report from the experts on the proposal in question (the Evaluation Summary Report).

International Cooperation Partner Countries (ICPC)

A list of low-income, lower-middle income and upper-middle-income countries, given in annex 1 to the work programme. Organisations from these countries can participate and receive funding in FP7, providing that certain minimum conditions are met.

International European Interest Organisation

International organisations, the majority of whose members are European Union Member states or Associated countries, and whose principal objective is to promote scientific and technological co-operation in Europe.

## **J**

Joint Research Centre (JRC)

The Commission's own research institutes.

## **L**

LEAR (Legal Entity Authorised Representative)

The LEAR is a person nominated in each legal entity participating in FP7. This person is the contact for the Commission related to all questions on legal status. He/she has access to the online database of legal entities with a possibility to view the data stored on his/her entity and to initiate updates and corrections to these data. The LEAR receives a Participant Identification Code (PIC) from the Commission (see below), and distributes this number within his/her organisation.

Lump sum

Lump sums do not require the submission of financial justifications (statements), as they are "fixed". ICPC participants when participating in an FP7 grant agreement have the choice between being reimbursed on the basis of eligible costs or on the basis of lump-sums. This choice can be made up to the moment of the signature of the grant agreement (whatever the final option chosen, the maximum EU contribution for the project remains unchanged). Once made, it will apply during the whole duration of the agreement without the possibility of changing it. ICPC participants may opt for a lump sum in a given project and for reimbursement of costs in another.

## **M**

Milestones

Control points where decisions are needed with regard to the next stage of the project.

## **N**

National Contact Points (NCP)

Official representatives nominated by the national authorities to provide tailored information and advice on each theme of FP7, in the national language(s).

Negotiation

The process of establishing a grant agreement between the Commission and an applicant whose proposal has been favourably evaluated, and when funds are available.

Non-profit

A legal entity is qualified as "non-profit" when considered as such by national or international law.

## **P**

### Part A

The part of a proposal dealing with administrative data. This part is completed using the web-based Participant Portal Submission Service.

### Part B

The part of a proposal explaining the work to be carried out, and the roles and aptitudes of the participants in the consortium. This part is uploaded to the Participant Portal as a pdf file

### Part B template

A document in PDF format supplied by the Participant Portal Submission Service, consisting of a template of all chapter headings, forms and tables required to prepare a proposal Part B. The template format is illustrated in Annex 4 to this Guide.

### Participants

The members of a consortium in a proposal or project. These are legal entities, and have rights and obligations with regard to the Community.

### Participant Identification Code (PIC)

Organisations participating in FP7 will progressively be assigned Participant Identification Codes (PIC). Possession of a PIC will enable organisations to take advantage of the Unique Registration Facility (see below), and to identify themselves in all transactions related to FP7 proposals and grants.

### Participant Portal

The single entry point for interaction with the research Directorates-General of the European Commission. It hosts a full range of services that facilitate the monitoring and the management of proposals and projects throughout their lifecycle, including calls for proposals, and access to the Participant Portal Submission Service.

### Participant Portal Submission Service

A web-based service, which must be used to submit proposals to the Commission. Access is given through the Participant Portal.

### Participant Portal Submission Service Helpdesk

A telephone / email service to assist applicants who have difficulty in submitting their proposal via the Participant Portal Submission Service :

tel: +32 2 29 92222 email [DIGIT-EFP7-SEP-SUPPORT@ec.europa.eu](mailto:DIGIT-EFP7-SEP-SUPPORT@ec.europa.eu)

### Programme committee

A group of official national representatives who assist the Commission in implementing the Framework Programme.

### Proposal

A description of the planned research activities, information on who will carry them out, how much they will cost, and how much funding is requested

### Public body

Public body means any legal entity established as such by national law, and international organisations.

## R

### Redress procedure

The initial information letter will indicate an address if an applicant wishes to submit a request for redress, if he or she believes that there have been shortcomings in the handling of the proposal in question, and that these shortcomings would jeopardise the outcome of the evaluation process. An internal evaluation review committee ("redress committee") will examine all such complaints. This committee does not itself evaluate the proposal. It is possible that the committee will recommend a re-evaluation of all or part of the proposal.

### Research organisation

A legal entity established as a non-profit organisation which carries out research or technological development as one of its main objectives.

### Reserve list

Due to budgetary constraints it may not be possible to support all proposals that have been evaluated positively. In such conditions, proposals on a reserve list will only be financed if funds become available following the negotiation of proposals on the main list.

### Risk-Sharing Finance Facility (RSFF)

A new mechanism to foster private sector investment in research by increasing the capacity of the EIB and its financial partners to provide loans for European RTD projects.

### RTD

Research and Technological Development.

## S

### SME

SMEs are micro, small and medium-sized enterprises. SMEs are defined in Recommendation 2003/361/EC of 6 May 2003.

### Specific flat rate (60%)

A 60% flat rate of the total direct costs applicable under certain conditions to non-profit public bodies, secondary and higher education establishments, research organisations and SMEs. This rate is available for the entire duration of FP7.

### Specific International Cooperation Actions (SICA)

In some calls on topics of mutual interest, special conditions apply to promote research collaborations between European organisations and those based in the International Cooperation Partner Countries (ICPC). This usually entails a minimum of two participants from EU or Associated countries, and two from ICPC.

## T

### Thresholds

For a proposal to be considered for funding, the evaluation scores for individual criteria must exceed certain thresholds. There is also an overall threshold for the sum of the scores.

### Two-stage submission

Some calls require proposals to be submitted in two stages. In this case, applicants initially present their idea in a brief outline proposal. This is evaluated against evaluation criteria, or sub-criteria for this stage set out in the call. Applicants successful in the first stage will be invited to submit a full proposal at the second stage, which will be evaluated against criteria for this second stage set out in the call. The first stage criteria, as set out in the work programme, are usually a limited set of those applying at the second stage.

#### Two-step evaluation

An evaluation procedure in which a proposal is evaluated first on a limited number of evaluation criteria (usually, just one), and only those proposals which achieve the threshold on this are subject to a full evaluation on the remaining criteria.

## **U**

### Unique Registration Facility (URF)

A system that will allow organisations who intend to submit on several occasions to register their details once and for all, obviating the need to provide the same information with each submission. The Web interface of the URF is found at <http://ec.europa.eu/research/participants/urf>. On this website you will also find a search tool to check if your organisation is already registered or not.

## **W**

### Weightings

The scores for certain evaluation criteria may be multiplied by a weighting factor before the total score is calculated. Generally, weightings are set to one; but there may be exceptions and applicants should check the details in annex 2 to this Guide.

### Work Package

A work package is a major sub-division of the proposed project with a verifiable end-point – normally a deliverable or a milestone in the overall project.

### Work Programme

A formal document of the Commission for the implementation of a specific programme, that sets out the research objectives and topics to be addressed. It also contains information that is set out further in this Guide, including the schedule and details of the calls for proposals, indicative budgets, and the evaluation procedure.

## ***Annexes***

- Annex 1 Timetable and specific information for this call
- Annex 2 Evaluation criteria and procedure
- Annex 3 Instructions for completing Part A of the proposal
- Annex 4 Instructions for drafting Part B of the proposal
- Annex 5 Ethical Guidelines for undertaking ICT research in FP7
- Annex 6 Pre-proposal check form



## *Annex 1: Timetable and specific information for this call*

- The **ICT part of the Cooperation work programme** provides the essential information for submitting a proposal to this call. It describes the content of the topics to be addressed, and details on how it will be implemented. The work programme is available on the CORDIS call page. You must consult this document.
- **Indicative timetable**

ICT Call 10 closes at 17h Brussels time on 15<sup>th</sup> January 2013

Publication of call	10 <sup>th</sup> July 2012
Deadline for submission of proposals	15 <sup>th</sup> January 2013 17h00 Brussels time
Evaluation of proposals	February-March 2013
Evaluation Summary Reports sent to all proposal coordinators	Mid April 2013
Invitation letter to successful applicants to launch negotiations with Commission services	End April 2013
Letters to unsuccessful applicants	May-June 2013
Signature of first grant agreements	July 2013

- **Further information and help**

### **Call information**

Participant Portal

<http://ec.europa.eu/research/participants/portal/>

### **General sources of help**

National Contact Points

[http://cordis.europa.eu/fp7/ncp\\_en.html](http://cordis.europa.eu/fp7/ncp_en.html)

FP7 Research enquiries service

<http://ec.europa.eu/research/enquiries>

Participant Portal Submission Service

Help desk tel: +32 2 299 2222

email [DIGIT-EFP7-SEP-SUPPORT@ec.europa.eu](mailto:DIGIT-EFP7-SEP-SUPPORT@ec.europa.eu)

Ethics Help desk

[http://cordis.europa.eu/fp7/get-support\\_en.html](http://cordis.europa.eu/fp7/get-support_en.html)

(information under "Ethics Help Desk for all FP7 projects")

Risk sharing financing facility  
(European Investment Bank)

<http://www.eib.org/rsff>

### **FP7/ICT Support projects**

Idealist partner search project

<http://www.ideal-ist.eu/>

IPR helpdesk

<http://www.ipr-helpdesk.eu/>

### **Legal documents generally applicable**

Decision on the Framework Programme  
Rules for Participation  
Specific Programmes  
Rules for proposal submission, evaluation selection and award

### **Contractual information**

Consortium agreement checklist  
Negotiation guidance notes  
Financial guidelines  
Model Grant agreement

All the above at [http://cordis.europa.eu/fp7/find-doc\\_en.html](http://cordis.europa.eu/fp7/find-doc_en.html)

- **Pre-proposal check**

The Commission offers a facility to allow a proposer to check on the appropriateness of their proposed action and the eligibility of the proposal consortium.

A form to request this check on your proposal is supplied as **Annex 6** of this Guide. This may be submitted at any time up to four weeks before the close of call, but it is wisest of course make this check as early as you can in your proposal preparation process.

The advice given by the Commission is strictly informal and non-binding. Only one pre-proposal check will be carried out per proposal. The advice provided through the pre-proposal check does not in any way engage the Commission with respect to acceptance or rejection of the proposal when it is formally submitted at a later stage. The evaluators who later evaluate your proposal will not be informed of the results of the pre-proposal check, nor even that a pre-proposal check was carried out. The pre-proposal service is not intended to assist with the identification of possible partners for your consortium.

Although this pre-proposal assessment service is entirely optional it is highly recommended to use this facility. Any proposal can always be submitted directly to the call without a pre-proposal check.

- **Address for pre-proposal check**

Please email your pre-proposal check form for this call to the address given on the "Contacts" document available on the call page on the Participant Portal.

## ***Annex 2: Evaluation criteria and procedures to be applied to CP-CSA for PCP proposals in this call***

### **1. General**

All eligible proposals will be evaluated by independent experts.

- Commission staff ensures that the process is fair, and in line with the principles contained in the Commission's rules<sup>15</sup>.
- Experts perform evaluations on a personal basis, not as representatives of their employer, their country or any other entity. They are expected to be independent, impartial and objective, and to behave throughout in a professional manner. They sign an appointment letter, including a confidentiality and conflict of interest declaration before beginning their work. Confidentiality rules must be adhered to at all times, before, during and after the evaluation.

In addition, an independent expert or experts will be appointed by the Commission to observe the evaluation process from the point of view of its working and execution. The role of the observer(s) is to give independent advice to the Commission on the conduct and fairness of the evaluation sessions, on the way in which the experts apply the evaluation criteria, and on ways in which the procedures could be improved. The observer(s) will not express views on the proposals under examination or the experts' opinions on the proposals.

### **2. Before the evaluation**

On receipt by the Commission, proposals are registered and acknowledged and their contents entered into a database to support the evaluation process. Eligibility criteria for each proposal are also checked by Commission staff before the evaluation begins. Proposals which do not fulfil these criteria will not be included in the evaluation.

For this call a proposal will only be considered eligible if it meets all of the following conditions:

- It is received by the Commission via the Participant Portal Submission Service before the deadline given in the call fiche
- It involves at least the minimum number of participants given in the call fiche
- It is complete (i.e. both the requested administrative forms and the proposal description are present)
- The content of the proposal relates to the topics and funding schemes, including any special conditions, set out in the relevant parts of the work programme

A proposal in which the Part B pdf file has been password-protected or for which printing has been blocked will be considered as failing the eligibility criteria under the third bullet point.

The Commission establishes a list of experts capable of evaluating the proposals that have been received. The list is drawn up to ensure:

- A high level of expertise;
- An appropriate range of competencies;

Provided that the above conditions can be satisfied, other factors are also taken into consideration:

- An appropriate balance between academic and industrial expertise and users;
- A reasonable gender balance;
- A reasonable distribution of geographical origins;

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<sup>15</sup> Rules on submission of proposals, and the related, evaluation, selection and award procedures [available on CORDIS]

- Regular rotation of experts

In constituting the lists of experts, the Commission also takes account of their abilities to appreciate the industrial and/or societal dimension of the proposed work. Experts must also have the appropriate language skills required for the proposals to be evaluated.

Commission staff allocates proposals to individual experts, taking account of the fields of expertise of the experts, and avoiding conflicts of interest.

### 3. Evaluation of proposals

At the beginning of the evaluation, experts will be briefed on the evaluation procedure, the experts' responsibilities, the issues involved in the particular area/objective, and other relevant material. The proposal will be evaluated against pre-determined evaluation criteria.

For CP-CSAs (combination of CP and CSA), according to Annex 2 of the work programme, criteria for "all funding schemes", "collaborative projects" and "coordination and support actions" apply. For a full understanding, please read also carefully Annex 4 of this Guide and Appendix 6 of the work programme.

<p align="center"><b>Evaluation criteria applicable to Pre-commercial Procurement (PCP) project proposals (CP-CSA)</b></p>		
<p><b>S/T QUALITY</b></p> <p><b>“Scientific and/or technological excellence (relevant to the topics addressed by the call)”</b></p>	<p><b>IMPLEMENTATION</b></p> <p><b>“Quality and efficiency of the implementation and the management”</b></p>	<p><b>IMPACT</b></p> <p><b>“Potential impact through the development, dissemination and use of project results”</b></p>
<ul style="list-style-type: none"> <li>• Soundness of concept and quality of objectives.</li> <li>• Progress beyond the state-of-the-art (relevant only to CP part of the proposal).</li> <li>• Contribution to the coordination of high quality research (relevant only to CSA part of the proposal).</li> <li>• Quality and effectiveness of the CSA mechanisms (mechanisms proposed to achieve the objectives of the networking and coordination CSA part of the project), and associated work plan</li> <li>• Quality and effectiveness of the S/T methodology and associated work plan (relevant only to CP part of the proposal).</li> </ul>	<ul style="list-style-type: none"> <li>• Appropriateness of the management structure and procedures.</li> <li>• Quality and relevant experience of the individual participants.</li> <li>• Quality of the consortium as a whole (including complementarity, balance).</li> <li>• Appropriate allocation and justification of the resources to be committed (staff, equipment ...).</li> </ul>	<ul style="list-style-type: none"> <li>• Contribution at the European level, to the expected impacts listed in the work programme under relevant topic/activity.</li> <li>• Appropriateness of measures for the exploitation of project results, dissemination of knowledge, through the engagement with stakeholders and the public at large, and the management of intellectual property and for spreading excellence.</li> </ul>

Evaluation scores will be awarded for each of the three criteria, not for the sub-criteria (bullet points). These sub-criteria are issues which the expert should consider in the assessment of that criterion. They also act as reminders of issues to raise later during the discussions of the proposal.

The relevance of a proposal will be considered in relation to the topic(s) of the work programme open in a given call, and to the objectives of a call. These aspects will be integrated in the application of the criterion "S/T quality", and the first sub-criterion under "Impact" respectively. When a proposal is partially relevant because it only marginally addresses the topic(s) of the call, or if only part of the proposal addresses the topic(s), this condition will be reflected in the scoring of the first criterion. Proposals that are clearly not relevant to a call ("out of scope") will be rejected on eligibility grounds.

Each criterion will be scored out of 5. Half marks can be given.

The scores indicate the following with respect to the criterion under examination:

- |     |   |
|-----|---|
| 0 - | <i>The proposal fails to address the criterion under examination or cannot be judged due to missing or incomplete information</i>           |
| 1 - | <b>Poor.</b> <i>The criterion is addressed in an inadequate manner, or there are serious inherent weaknesses.</i>                           |
| 2 - | <b>Fair.</b> <i>While the proposal broadly addresses the criterion, there are significant weaknesses.</i>                                   |
| 3 - | <b>Good.</b> <i>The proposal addresses the criterion well, although improvements would be necessary.</i>                                    |
| 4 - | <b>Very good.</b> <i>The proposal addresses the criterion very well, although certain improvements are still possible.</i>                  |
| 5 - | <b>Excellent.</b> <i>The proposal successfully addresses all relevant aspects of the criterion in question. Any shortcomings are minor.</i> |

No weightings will be applied.

Thresholds will be applied to the scores. The threshold for individual criteria will be 3. The overall threshold, applying to the sum of the three individual scores, will be 10.

Conflicts of interest: Under the terms of their appointment letter, experts must declare beforehand any known conflicts of interest, and must immediately inform a Commission staff member if one becomes apparent during the course of the evaluation. The Commission will take whatever action is necessary to remove any conflict.

Confidentiality: The appointment letter also requires experts to maintain strict confidentiality with respect to the whole evaluation process. They must follow any instruction given by the Commission to ensure this. Under no circumstance may an expert attempt to contact an applicant on his own account, either during the evaluation or afterwards.

#### **4. Individual evaluation**

The first stage (individual evaluation) will be carried out on the premises of the experts concerned ("remotely").

Each proposal will first be assessed independently by five or more experts, chosen by the Commission from the pool of experts taking part in this evaluation. At this first step the experts are acting individually; they do not discuss the proposal with each other, nor with any third party. The experts record their individual opinions in an **Individual Evaluation Report (IER)**, giving scores and also comments against the evaluation criteria.

When scoring proposals, experts must *only* apply the above evaluation criteria.

Experts will assess and mark the proposal exactly as it is described and presented. They do not make any assumptions or interpretations about the project in addition to what is in the proposal.

Concise but explicit justifications will be given for each score. Recommendations for improvements to be discussed as part of a possible negotiation phase will be given, if needed.

The experts will also indicate whether, in their view, the proposal deals with sensitive ethical issues.

Signature of the IER also entails a declaration that the expert has no conflict of interest in evaluating the particular proposal.

Scope of the call: It is possible that a proposal is found to be out of scope of the call during the course of the individual evaluation, and therefore not relevant. If an expert suspects that this may be the case, a Commission staff member will be informed immediately, and the views of the other experts will be sought. If the general view is that the main part of the proposal is not relevant to the topics of the call, the proposal will be withdrawn from the evaluation, and the proposal will be deemed ineligible.

## 5. Consensus meeting

Once all the experts to whom a proposal has been assigned have completed their IER, the evaluation progresses to a consensus assessment, representing their common views.

This entails a consensus meeting to discuss the scores awarded and to prepare comments.

The consensus discussion is moderated by a representative of the Commission. The role of the moderator is to seek to arrive at a consensus between the individual views of experts without any prejudice for or against particular proposals or the organisations involved, and to ensure a confidential, fair and equitable evaluation of each proposal according to the required evaluation criteria.

The moderator for the group may designate an expert to be responsible for drafting the consensus report ("rapporteur"). The experts attempt to agree on a consensus score for each of the criteria that have been evaluated and suitable comments to justify the scores. Comments should be suitable for feedback to the proposal coordinator. Scores and comments are set out in a consensus report. They also come to a common view on the questions of scope if necessary.

The consensus group will also suggest questions to be asked during the hearing, if one is foreseen (see below).

If during the consensus discussion it is found to be impossible to bring all the experts to a common point of view on any particular aspect of the proposal, the Commission may ask up to three additional experts to examine the proposal.

Outcome of consensus: The outcome of the consensus step is the **Consensus Report (CR)**. This will be signed (either on paper, or electronically) by all experts, or as a minimum, by the rapporteur and the moderator. The moderator is responsible for ensuring that the consensus report reflects the consensus reached, expressed in scores and comments. In the case that it is impossible to reach a consensus, the report sets out the majority view of the experts but also records any dissenting views.

Ethical issues (above threshold proposals): If one<sup>16</sup> or more experts have noted that there are ethical issues touched on by the proposal, and the proposal is considered to be above threshold, the relevant box on the consensus report (CR) will be ticked and an Ethical Issues Report (EIR) completed, stating the nature of the ethical issues. The EIR will be signed by the Commission moderator and one member of the consensus group (normally, the proposal rapporteur).

The Commission will take the necessary steps to assure the quality of the consensus reports, with particular attention given to clarity, consistency, and appropriate level of detail. If important changes are necessary, the reports will be referred back to the experts concerned.

The signing of the consensus report completes the consensus step.

Evaluation of a resubmitted proposal: In the case of proposals that have been submitted previously to the Commission in FP7, the moderator may give the experts the previous evaluation summary report (see below) following the consensus stage. If necessary, the experts will be required to provide a clear justification for their scores and comments should these differ markedly from those awarded to the earlier proposal.

## 6. Panel review

This is the final step involving the independent experts. It allows them to formulate their recommendations to the Commission having had an overview of the results of the consensus step.

The panel comprises experts involved at the consensus step with the experts who reviewed the other proposals in the area.

The main task of the panel is to examine and compare the consensus reports in a given area, to check on the consistency of the marks applied during the consensus discussions and, where necessary, propose a new set of consensus scores.

The tasks of the panel will also include:

- resolving cases where a minority view was recorded in the consensus report
- recommending a priority order for proposals with the same score
- making recommendations on possible clustering or combination of proposals.

The panel is chaired by the Commission. The Commission will ensure fair and equal treatment of the proposals in the panel discussions. A panel rapporteur will be appointed to draft the panel's advice. A ranked list will be drawn up for every indicative budget as shown in the call fiche. The panel can deal with one or more ranked lists for the proposals under evaluation, following the scoring systems indicated above.

### Priority order for proposals with the same score

As part of the evaluation by independent experts, a panel review will recommend one or more ranked lists for the proposals under evaluation, following the scoring systems indicated above. A ranked list will be drawn up for every indicative budget shown in the call fiche.

If necessary, the panel will determine a priority order for proposals which have been awarded the same score within a ranked list. Whether or not such a prioritisation is carried out will depend on the available budget or other conditions set out in the call fiche. The following approach will be applied successively for every group of *ex aequo* proposals requiring prioritisation, starting with the highest scored group, and continuing in descending order:

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<sup>16</sup> Exceptionally for this issue, no consensus is required.

(i) Proposals that address relevant topics not otherwise covered by more highly-rated proposals will be considered to have the highest priority.

(ii) These proposals will themselves be prioritised according to the scores they have been awarded for the criterion *impact*. When these scores are equal, priority will be based on scores for the criterion *scientific and/or technological excellence*. If necessary, any further prioritisation will be based on other appropriate characteristics, to be decided by the panel, related to the contribution of the proposal to the European Research Area and/or general objectives mentioned in the work programme (e.g. presence of SMEs, international co-operation, public engagement).

(iii) The method described in (ii) will then be applied to the remaining *ex aequo* in the group.

The outcome of the panel meeting is a report recording, principally:

- An **Evaluation Summary Report** (ESR) for each proposal, including, where relevant, a report of any ethical issues raised;
- A list of proposals passing all thresholds, along with a final score for each proposal passing the thresholds and the panel recommendations for priority order;
- A list of evaluated proposals having failed one or more thresholds;
- A list of any proposals having been found ineligible;
- A summary of the deliberations of the panel.

If the panel has considered proposals submitted to various parts of a call (e.g. different funding schemes, or different topics that have been allocated distinct indicative budgets in the work programme), the report may accordingly contain multiple priority lists.

The panel report is signed by at least three panel experts and the Commission chairperson.

A copy of the Evaluation Summary Report will be sent to each proposal coordinator.

## 7. Ethics Review of project proposals

An ethics review of above-threshold proposals may be organised by the Commission. The Ethics Review is carried out by independent experts with a special expertise on ethics. Reviewing research projects on ethical grounds at the EU level is a legal requirement under FP7. The Review evaluates aspects of the design and methodology of the proposed research such as intervention on humans, use of animals, data protection issues, terms of participation of children and vulnerable populations groups.

The Panel drafts an Ethics Review Report that summarises its opinion on the ethical soundness of the project proposal under consideration. The requirements put forward by the Panel are taken into account in any subsequent negotiations on the grant agreement, and may lead to obligatory provisions in the conduct of the research.

For additional information on the Ethics Review procedure see:

<http://ec.europa.eu/research/science-society/index.cfm?fuseaction=public.topic&id=73>



### ***Annex 3: Instructions for completing Part A of the proposal***

Proposals in this call must be submitted electronically, using the Commission's Participant Portal Submission Service. The procedure is summarised in section 3 of this guide.

In Part A you are asked for certain administrative details that will be used in the evaluation and further processing of your proposal. Section A1 gives a snapshot of your proposal, section A2 concerns the participants in the consortium, while section A3 deals with money matters. Details of the work you intend to carry out will be described in Part B (see annex 4 of this guide).

Please note:

- The coordinator fills in the sections A1 and A3.
- Participants already identified at the time of proposal submission (including the coordinator) each fill in their respective section A2.
- Subcontractors, if any, do not fill in a section A2 and are not listed separately in section A3 (They are described in Part B)
- The estimated budget planned for any future participants, not yet identified at the time of the proposal, is not shown separately in form A3 but must be added to the coordinator's budget. (Their role, profile and tasks are described in Part B)

When you complete section A3, please make sure that:

- *Numbers are always rounded to the nearest whole number*
- *You have inserted zeros ("0") where there are no costs or funding figures. Leaving cells empty will block the submission of your proposal*
- *All costs are given in Euros (not thousands of Euros)*
- *Costs do not include Value Added Tax*

The following notes are for information only. They should assist you in completing the A-part of your proposal. On-line guidance will also be available. The precise questions and options presented on the Participant Portal Submission Service may differ slightly from these below.

<b>Section A1: Summary</b>	
<b>Proposal Acronym</b>	<p>The short title or acronym will be used to identify your proposal efficiently in this call. It should be of <u>no more than 20 characters</u> (use standard alphabet and numbers only; no symbols or special characters please).</p> <p>The same acronym should appear on each page of part B of your proposal.</p>
<b>Collaborative Projects &amp; Coordination and Support Actions</b>	<p>For each type of Collaborative Projects and Coordination and Support Actions, please refer to the work programme.</p>
<b>Proposal Title</b>	<p>The title should be <u>no longer than 200 characters</u> and should be understandable to the non-specialist in your field.</p>
<b>Duration in months</b>	<p>Insert the estimated duration of the project in full months.</p>
<b>Call (part) identifier</b>	<p>The call identifier is the reference number given in the call or part of the call you are addressing, as indicated in the publication of the call in the Official Journal of the European Union, and on the CORDIS call page.  <b>[The call identifier is pre-filled in the forms from the Participant Portal Submission Service. If you do not have the correct identifier on your forms, you have registered for the wrong call. Discard this registration and register again].</b></p>
<b>Topic code(s) most relevant to your proposal</b>	<p>All activities and topics of FP7 have been assigned unique codes, which are used in the processing of data on proposals and subsequent contracts. The codes are organised hierarchically.</p> <p>The choice of the first activity code will be limited in the drop-down menu to one of the topics open in this call. Select the code corresponding to the topic most relevant to your proposal.</p> <p>The choice for the second code is also limited to topics open in the call in question. Enter a second code if your proposal also addresses another of these. Select 'none' if this is not the case.</p> <p>Select a third code if your proposal is also relevant to another theme. This time, the available codes will simply correspond to broad themes. Select 'none' if this is not the case.</p>
<b>Free Keywords</b>	<p>Please enter a number of keywords that you consider sufficient to characterise the scope of your proposal.</p> <p>There is <u>a limit of 100 characters</u>.</p>
<b>Abstract</b>	<p>The abstract should, at a glance, 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. Please use plain typed text, avoiding formulae and other special characters. If the proposal is written in a language other than English, please include an English version of the proposal abstract in Part B.</p> <p><b>There is a limit of 2000 characters. Exceeding this limit may block the submission of your proposal!</b></p>
<b>Similar proposals or signed contracts</b>	<p>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.</p>

<b>Section A2/ Participants</b>	
<b>Participant number</b>	The number allocated by the consortium to the participant for this proposal. The <b>co-ordinator</b> of a proposal is always <b>number one</b> .
<b>Participant Identify Code</b>	The Participant Identification Code (PIC) enables organisations to take advantage of the Participant Portal. Organisations who have received a PIC from the Commission are encouraged to use it when submitting proposals. By entering a PIC, parts of section A2 will be filled in automatically. An online tool to search for existing PICs and the related organisations is available at <a href="http://ec.europa.eu/research/participants/portal/page/searchorganisations">http://ec.europa.eu/research/participants/portal/page/searchorganisations</a> . Organisations not yet having a PIC are strongly encouraged to register before submitting the proposal and insert in section A2 the temporary PIC received at the end of the registration process..
<b>Legal name</b>	<p><b>For Public Law Body</b>, it is the name under which your organisation is registered in the Resolution text, Law, Decree/Decision establishing the Public Entity, or in any other document established at the constitution of the Public Law Body;</p> <p><b>For Private Law Body</b>, it is the name under which your organisation is registered in the national Official Journal (or equivalent) or in the national company register.</p> <p><b>For a natural person</b>, it is for e.g. Mr Adam JOHNSON, Mrs Anna KUZARA, and Ms Alicia DUPONT.</p>
<b>Organisation Short Name</b>	<p>Choose an abbreviation of your Organisation Legal Name, only for use in this proposal and in all relating documents.</p> <p>This short name should not be more <u>than 20 characters</u> exclusive of special characters (./;...), for e.g. CNRS and not C.N.R.S. It should be preferably the one as commonly used, for e.g. IBM and not Int.Bus.Mac.</p>
<b>Legal address</b>	<p>For Public and Private Law Bodies, it is the address of the entity's Head Office.</p> <p>For Natural persons it is the Official Address.</p> <p>If your address is specified by an indicator of location other than a street name and number, please insert this instead under the "street name" field and "N/A" under the "number" field.</p>
<b>Non-profit organisation</b>	Non-profit organisation is a legal entity qualified as such when it is recognised by national or, international law.
<b>Public body</b>	Public body means any legal entity established as such by national law, and international organisations.
<b>Research organisation</b>	Research organisation means a legal entity established as a non-profit organisation which carries out research or technological development as one of its main objectives.
<b>NACE code</b>	<p><b>NACE</b> means "<u>N</u>omenclature des <u>A</u>ctivités économiques dans la <u>C</u>ommunauté <u>E</u>uropéenne".</p> <p>Please select <b>one</b> activity from the list that <b>best</b> describes your professional and economic ventures. If you are involved in more than one economic activity, please select the <b>one</b> activity that is <b>most</b> relevant in the context of your contribution to the proposed project. For more information on the methodology, structure and full content of NACE (rev. 1.1) classification please consult EUROSTAT at:</p> <p><a href="http://ec.europa.eu/eurostat/ramon/nomenclatures/index.cfm?TargetUrl=LST_CLS_DLD&amp;StrNom=NACE_1_1&amp;StrLanguageCode=EN&amp;StrLayoutCode=HIERARCHIC">http://ec.europa.eu/eurostat/ramon/nomenclatures/index.cfm?TargetUrl=LST_CLS_DLD&amp;StrNom=NACE_1_1&amp;StrLanguageCode=EN&amp;StrLayoutCode=HIERARCHIC</a> .</p>

<p><b>Small and Medium-Sized Enterprises (SMEs)</b></p>	<p>SMEs are micro, small and medium-sized enterprises within the meaning of Recommendation 2003/361/EC in the version of 6 May 2003. The full definition and a guidance booklet can be found at <a href="http://ec.europa.eu/enterprise/enterprise_policy/sme_definition/index_en.htm">http://ec.europa.eu/enterprise/enterprise_policy/sme_definition/index_en.htm</a></p> <p>To find out if your organisation corresponds to the definition of an SME you can use the on-line tool at <a href="http://ec.europa.eu/research/sme-techweb/index_en.cfm">http://ec.europa.eu/research/sme-techweb/index_en.cfm</a></p>
<p><b>Dependencies with (an) other participant(s)</b></p>	<p>Two participants (legal entities) are dependent on each other where there is a controlling relationship between them:</p> <ul style="list-style-type: none"> <li>– A legal entity is under the same direct or indirect control as another legal entity (<b>SG</b>);</li> <li>or</li> <li>– A legal entity directly or indirectly controls another legal entity (<b>CLS</b>);</li> <li>or</li> <li>– A legal entity is directly or indirectly controlled by another legal entity (<b>CLB</b>).</li> </ul> <p><b>Control:</b> Legal entity A controls legal entity B if:</p> <ul style="list-style-type: none"> <li>– 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,</li> <li>or</li> <li>– A, directly or indirectly, holds in fact or in law the decision-making powers in B.</li> </ul> <p>The following relationships between legal entities shall not in themselves be deemed to constitute controlling relationships:</p> <ul style="list-style-type: none"> <li>(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;</li> <li>(b) the legal entities concerned are owned or supervised by the same public body.</li> </ul>
<p><b>Character of dependence</b></p>	<p>According to the explanation above mentioned, please insert the appropriate abbreviation according to the list below to characterise the relation between your organisation and the other participant(s) you are related with:</p> <ul style="list-style-type: none"> <li>• <b>SG</b>: Same group: if your organisation and the other participant are controlled by the same third party;</li> <li>• <b>CLS</b>: Controls: if your organisation controls the other participant;</li> <li>• <b>CLB</b>: Controlled by: if your organisation is controlled by the other participant.</li> </ul>
<p><b>Contact point</b></p>	<p>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 Commission will contact concerning this proposal (e.g. for additional information, invitation to hearings, sending of evaluation results, convocation to negotiations).</p>
<p><b>Title</b></p>	<p>Please choose one of the following: Prof., Dr., Mr., Mrs, Ms.</p>
<p><b>Sex</b></p>	<p>This information is required for statistical and mailing purposes. Indicate F or M as appropriate.</p>
<p><b>Phone and fax numbers</b></p>	<p>Please insert the full numbers including country and city/area code. Example +32-2-2991111.</p>
<p><b>Section A3/Budget</b></p>	
<p><b>Indirect Costs</b></p>	<p>Indirect costs are all those eligible costs which cannot be identified by the participant as being directly attributed to the project but which can be identified and justified by its accounting system as being incurred in direct relationship with the eligible direct costs attributed to the project. They may not include any eligible direct costs.</p>

<b>Method of calculating indirect costs*</b>	<p><b>Summary description</b></p> <p>Participants who have an analytical accounting system that can identify and group their indirect costs in accordance with the eligibility criteria (e.g. exclude non-eligible costs) must report their actual indirect costs (or choose the 20% flat rate option referred to below).</p> <p>For the purpose of calculating the actual indirect costs, a participant is allowed to use a simplified method of calculation of its full indirect eligible costs.</p> <p>Optionally, participants may opt for a flat rate for indirect costs of 20% of the direct costs (minus subcontracting and third party costs not incurred on the premises of the participant).</p> <p>A specific flat rate of 60% of the direct costs is foreseen for non-profit public bodies, secondary and higher education establishments, research organisations and SMEs which are unable to identify with certainty their real indirect costs for the project.*</p> <p>For <b>Coordination and Support actions</b>, whichever method is used, the reimbursement of indirect eligible costs may not exceed 7% of the direct eligible costs, excluding the direct eligible costs for subcontracting and the costs of reimbursement of resources made available by third parties which are not used on the premises of the participant.</p> <p><b>Further guidance*</b></p> <p>In FP7 all departments, faculties or institutes which are part of the same legal entity must use the same system of cost calculation (unless a special clause foreseeing a derogation for a particular department/institute is included in the grant agreement). Under FP7, there are no cost reporting models.</p> <p>1. Participants which have an analytical accounting system that can identify and group their indirect costs (pool of costs) in accordance with the eligibility criteria (e.g. exclude non-eligible costs) must report their actual indirect costs (or choose the 20% flat rate option under 2. below). This method is the same as the "full cost" model used in previous Framework Programmes.</p> <p>For the purpose of calculating the actual indirect costs, a participant is allowed to use a simplified method of calculation of its full indirect eligible costs. The simplified method is a way of declaring indirect costs which applies to organisations which do not aggregate their indirect costs at a detailed level (centre, department), but can aggregate their indirect costs at the level of the legal entity.</p> <p>The simplified method can be used if the organisation does not have an accounting system with a detailed cost allocation. The method has to be in accordance with their usual accounting and management principles and practices; it does not involve necessarily the introduction of a new method just for FP7 purposes. Participants are allowed to use it, provided this simplified approach is based on actual costs derived from the financial accounts of the last closed accounting year.</p> <p>There is no "standard model"; each legal entity will use its own system. The minimum requirements for it to be considered a simplified method for FP7 purposes are the following:</p> <ul style="list-style-type: none"> <li>- the system must allow the participant to identify and remove its direct ineligible costs (VAT, etc.);</li> <li>- it must at least allow for the allocation of the overheads at the level of the legal entity to the individual projects by using a fair "driver" (e.g. total productive hours);</li> <li>- the system applied and the costs declared according to it should follow the normal accounting principles and practices of the participant. Therefore, if the system used by a participant is more "refined" than the "minimum" requirements mentioned here, it is that system which should be used when declaring costs.</li> </ul> <p>Example: if a participant's accounting system distinguishes between different overheads rates according to the type of activity (research, teaching...), then the overheads declared in an FP7 grant agreement should follow this practice and refer only to the concerned activities (research, demonstration...)</p> <p>The simplified method does not require previous registration or certification by the Commission.</p> <p>2. Optionally, participants may opt to declare their actual direct costs plus a flat rate for indirect costs of 20% of the direct costs (minus subcontracting and third party costs not incurred on the premises of the participant). This flat rate is open to any participant whatever the accounting system it uses. Accordingly, when this option is chosen, there is no need for certification of the indirect costs, only of the direct ones.</p> <p>3. Also, a <b>specific flat rate</b> is foreseen for certain types of organisations.</p> <p>The use of this flat rate is subject to three cumulative conditions:</p> <p>(i) Status of the organisation The flat rate is reserved to:</p> <ul style="list-style-type: none"> <li>- non-profit public bodies</li> <li>- secondary and higher education establishments</li> </ul>
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	<p>- research organisations - SMEs</p> <p>(ii) Accounting system of the organisation The flat rate is foreseen for the organisations which are unable to identify with certainty their real indirect costs for the project. How will it be proved that an organisation is unable to identify with certainty their real indirect costs for the project? The participant (for example, an SME) does not have to change its accounting system or its usual accounting principles. If its accounting system can identify overall overheads but does not allocate them to project costs, then the participant can use this flat rate if the other conditions are fulfilled. Example: A University, which in FP6 has used the "additional cost" basis because its accounting system did not allow for the share of their direct and indirect costs to the project to be distinguished may under FP7: - either opt for the 60% flat rate, or - introduce a cost accounting system "simplified method" by which a basic allocation per project of the overhead costs of the legal entity will be established, or - introduce a full analytical accounting system.</p> <p>Following this, an organisation which used the "full cost" model under the Sixth Framework Programme is presumed to be in a situation to be able to identify the real indirect costs and allocate them to the projects. Accordingly, this organisation would not in principle be able to opt for the 60% flat rate for FP7.</p> <p>An organisation which can identify the real indirect costs but does not have a system to allocate these indirect costs can opt for this 60% flat rate. The choice of this specific flat rate lies within the responsibility of the participant. If a subsequent audit shows that the above-mentioned cumulative conditions are not fulfilled, all projects where this participant is involved might be reviewed.</p> <p>(iii) Type of funding scheme The flat rate is reserved to funding schemes which include research and technological development and demonstration activities: Network of Excellence and Collaborative projects (including research for the benefit of specific groups – in particular SMEs). The basis for the calculation of the flat rate excludes the costs for subcontracting and the costs of resources made available by third parties which are not used on the premises of the participant because in these two cases, the indirect costs are not incurred by the participant but by the subcontractor or the third party. When a participant opts for the specific flat rate of 60 % for its first participation under FP7 it can opt afterwards for the actual indirect costs system for subsequent participations. This change does not affect previous grant agreement. After this change, this organisation cannot opt again for a flat rate system (either 60% or 20% flat rate).</p> <p>(*) In conformity with the FP7 Rules of participation, for the CP part, the reimbursement is limited to maximum 75% of the eligible (direct) costs for the development of the new ICT solutions procured through the joint PCP call for tender. It does not cover indirect costs (overheads).</p>
<p><b>International Cooperation Partner Country (ICPC)</b></p>	<p>International Cooperation Partner Country means a third country which the Commission classifies as a low-income, lower-middle income or upper-middle-income country and which is identified as such in Annex I of the work programmes.</p>
<p><b>Lump sum funding method</b></p>	<p>Legal entities established in an ICPC may opt for lump sums. In that case the contribution is based on the amounts shown below, multiplied by the total number of person-years for the project requested by the ICPC legal entity.</p> <ul style="list-style-type: none"> <li>• Low-income ICPC: 8,000 Euro/researcher/year</li> <li>• Lower middle income ICPC: 9,800 Euro/researcher/year</li> <li>• Upper middle income ICPC 20,700 Euro/researcher/year</li> </ul> <p>The maximum EU contribution is calculated by applying the normal upper funding limits shown under "requested EU contribution". This amount is all inclusive, covering support towards both the direct and the indirect costs.</p> <p>More information on ICPC lump sums can be found in the section II.18 of the "Guide to financial issues" <a href="http://cordis.europa.eu/fp7/find-doc_en.html">http://cordis.europa.eu/fp7/find-doc_en.html</a></p>

<p><b>Type of Activity</b></p>	<ul style="list-style-type: none"> <li>• <b>RTD activities</b> are directly aimed at creating new knowledge and new technology. For PCP CP-CSAs it includes the costs of <b>joint research activities</b>*.</li> <li>• <b>Coordination activities</b> for the development and implementation of common, joint, strategic PCP activities. For PCP CP-CSAs it includes the costs of <b>networking and coordination activities</b>* (including, but not limited to, training, dissemination and communication). It does not cover management costs.</li> <li>• <b>Management activities</b> include the maintenance of the consortium agreement, if it is obligatory, the overall legal, ethical, financial and administrative management including for each of the participants obtaining the certificates on the financial statements or on the methodology, the implementation of competitive calls by the consortium for the participation of new participants and, any other management activities foreseen in the proposal* except coordination of research and technological development activities.</li> <li>• <b>Other activities</b> means any specific activities not covered by the above mentioned types of activity. These activities should be specified in the proposal Part B.</li> </ul> <p>(*) as defined in Section 2.2 of this Guide for Applicants.</p>
<p><b>Personnel costs</b></p>	<p>Personnel costs are only the costs of the actual hours worked by the persons directly carrying out work under the project and shall reflect the total remuneration: salaries plus social security charges (holiday pay, pension contribution, health insurance, etc.) and other statutory costs included in the remuneration. Such persons must:</p> <ul style="list-style-type: none"> <li>– be directly hired by the participant in accordance with its national legislation,</li> <li>– be working under the sole technical supervision and responsibility of the latter, and</li> <li>– be remunerated in accordance with the normal practices of the participant.</li> </ul> <p>Participants may opt to declare average personnel costs if certified in accordance with a methodology approved by the Commission and consistent with the management principles and usual accounting practices of the participant. Average personnel costs charged by a participant having provided a certification on the methodology are deemed not to significantly differ from actual personnel costs.</p>
<p><b>Sub-contracting</b></p>	<p>A subcontractor is a third party which has entered into an agreement on business conditions with one or more participants, in order to carry out part of the work of the project without the direct supervision of the participant and without a relationship of subordination.</p> <p>Where it is necessary for the participants to subcontract certain elements of the work to be carried out, the following conditions must be fulfilled:</p> <ul style="list-style-type: none"> <li>- subcontracts may only cover the execution of a limited part of the project*;</li> <li>- recourse to the award of subcontracts must be duly justified in Part B of the proposal having regard to the nature of the project and what is necessary for its implementation;</li> </ul> <p>recourse to the award of subcontract by a participant may not affect the rights and obligations of the participants regarding background and foreground**;</p> <ul style="list-style-type: none"> <li>- Part B of the proposal must indicate the task to be subcontracted and an estimation of the costs;</li> </ul> <p>Any subcontract, the costs of which are to be claimed as an eligible cost, must be awarded according to the principles of best value for money (best price-quality ratio), transparency and equal treatment. Framework contracts between a participant and a subcontractor, entered into prior to the beginning of the project that are according to the participant's usual management principles may also be accepted.</p> <p>Participants may use external support services for assistance with minor tasks that do not represent per se project tasks as identified in Part B of the proposal.</p> <p>If applicable, actual direct costs and real overhead costs of third parties that make available to the proposal resources otherwise unavailable within the consortium, can also be included under the category of subcontracting costs (provided that these costs are not related to proposal's core tasks).</p> <p>(*) In conformity with the work programme, for the CP-CSAs for PCP, the eligible costs for the development of ICT solutions that are procured by the participants from subcontractors selected through the joint PCP call for tender, can be reimbursed up to maximum 75% (75% for non-profit public bodies, secondary and higher education establishments, research organisations and SMEs which are unable to identify with certainty their real indirect costs for the project, 50% for other type of beneficiaries).</p> <p>(**) To this end, the CP-CSA for PCP is implemented according to the IPR conditions described in Appendix 6 of</p>

	the work programme
<b>Other direct costs</b>	Means direct costs not covered by the above mentioned categories of costs.
<b>Total Budget</b>	<b>[Note: The "total budget" is <u>not</u> the requested EU contribution]</b> A sum of all the eligible costs, under the respective types of activity.
<b>Requested EU contribution</b>	The requested EU contribution shall be determined by applying the upper funding limits indicated below, per activity and per participant to the costs accepted by the Commission, or to the flat rates or lump sums.  <b>Maximum reimbursement rates of eligible costs</b> <ul style="list-style-type: none"> <li>• <b>Research and technological development = 75%*</b></li> <li>• <b>Coordination activities = 100%</b></li> <li>• <b>Management activities = 100%</b></li> <li>• <b>Other activities = 100%</b></li> </ul> <p>(*) 75% for non-profit public bodies, secondary and higher education establishments, research organisations and SMEs which are unable to identify with certainty their real indirect costs for the project, 50% for other type of beneficiaries</p>
<b>Total Receipts</b>	<b>[Note: The term "receipts" is <u>not</u> the requested EU contribution.]</b>  Receipts of the project may arise from: <ul style="list-style-type: none"> <li>a) Financial transfers or contributions in kind free of charge to the participant from third parties: <ul style="list-style-type: none"> <li>i. shall be considered a receipt of the project if they have been contributed by the third party specifically to be used on the project.</li> <li>ii. shall <u>not</u> be considered a receipt of the project if their use is at the management discretion of the participant.</li> </ul> </li> <li>b) Income generated by the project: <ul style="list-style-type: none"> <li>i. shall be considered receipts for the participant when generated by actions undertaken in carrying out the project and from the sale of assets purchased under the grant agreement up to the value of the cost initially charged to the project by the participant;</li> <li>ii. shall <u>not</u> be considered a receipt for the participant when generated from the use of foreground resulting from the project.</li> </ul> </li> </ul> <p>The Community financial contribution may not have the purpose or effect of producing a profit for the participants. For this reason, the total requested EU funding plus receipts cannot exceed the total eligible costs.</p>



### *Annex 4: Instructions for drafting Part B of the proposal*

## **Combination of Collaborative Projects and Coordination and Support Actions (CP-CSA) for Pre-Commercial Procurement (PCP)**

A description of this funding scheme is given in section 2 of this Guide for Applicants. Please examine this carefully before preparing your proposal.

This Annex provides a template to help you structure your proposal. It will help you present important aspects of your planned work in a way that will enable the experts to make an effective assessment against the evaluation criteria (see Annex 2). Sections 1, 2 and 3 each correspond to an evaluation criterion. The sub-sections (1.1, 1.2 etc.) correspond to the sub-criteria.

Remember, please keep to the page limits where these are specified. Information given on excess pages may be disregarded. The minimum font size allowed is 11 points. All margins (top, bottom, left, right) should be at least 15 mm (not including any footers or headers).

Even where no page limits are given it is in your interest to keep your text concise, since over-long proposals are rarely viewed in a positive light by the experts.

### **Title Page**

Proposal full title

Proposal acronym

Type of funding scheme:

In this case - Combination of Collaborative Project and Coordination and Support Action: Pre-Commercial Procurement (PCP)

Work programme topic and subtopic addressed

(if more than one, indicate their order of importance to the project)

Name of the coordinating person

List of participants:

<b>Participant no. *</b>	<b>Participant organisation name</b>	<b>Part. short name</b>	<b>Country</b>
1 (Coordinator)			
2			
3			

\* Please use the same participant numbering as that used in Proposal submission forms A2

### **Proposal abstract**

(copied from Part A, if not in English include an English translation)

### **Table of Contents**

## Proposal

### **Section 1: Scientific and/or technical excellence, relevant to the topics addressed by the call**

#### **1.1 Soundness of concept and quality of objectives**

Explain the concept of your project. What are the main ideas that led you to propose this work? Describe in detail the S&T objectives. Show how they relate to the topics addressed by the call, which you should explicitly identify. The objectives should be those achievable within the project. They should be stated in a measurable and verifiable form, including through the milestones that will be indicated under section 1.4 and 1.5 below.

In particular, describe the overall approach presented in the proposal for combining networking and coordination activities with joint research activities (PCP procurement of new ICT developments) in view of bringing the targeted quality and efficiency improvements in the area of public interest addressed by the project.

#### **1.2 Progress beyond the state-of-the-art (relevant to CP part only)**

Describe the state-of-the-art in the area concerned, and the advance that the proposed project would bring about. In particular, describe the degree of ambition in the R&D services to be procured in the joint PCP (joint research activities – CP – part of the proposal). If applicable, refer to the results of any patent search you might have carried out.

In the proposal, consortia shall have jointly identified a concrete challenge in the mid-to-long term innovation plans of the participating public purchasers that requires new R&D, which is proposed to be procured in cooperation through PCP. The consortium shall provide justification in the proposal that the topic proposed for the joint PCP call for tender would fit the scope of an R&D<sup>17</sup> services contract<sup>18</sup>.

#### **1.3. Contribution to the co-ordination of high quality research (relevant to the CSA part only)**

Describe how the proposed joint networking and coordination activities strengthen the cooperation between public bodies in Europe in the innovation of their public services through a strategy that includes PCP, in particular through the preparation, management and coordination of a joint PCP call for tender.

#### **1.4 Coordination Activities (the CSA part of the project) and associated work plan**

Describe the extent to which the proposed co-ordination mechanisms will foster a culture of co-operation between the participants, and bring about the aspired innovations in public procurement strategies through PCP to enhance the quality and efficiency of public services to the citizens.

A detailed work plan should be presented, broken down into work packages<sup>19</sup> (WPs) which should follow the logical phases of the implementation of the project's Coordination and Networking

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<sup>17</sup> R&D can cover activities such as solution exploration and design, prototyping, up to the original development of a limited volume of first products or services in the form of a test series. Original development of a first product or service may include limited production or supply in order to incorporate the results of field testing and to demonstrate that the product or service is suitable for production or supply in quantity to acceptable quality standards. R&D does not include commercial development activities such as quantity production, supply to establish commercial viability or to recover R&D costs, integration, customisation, incremental adaptations and improvements to existing products or processes.

<sup>18</sup> Contracts providing more than only services are still considered a public service contract if the value of the services exceeds that of the products covered by the contract.

<sup>19</sup> A work package is a major sub-division of the proposed project with a verifiable end-point - normally a deliverable or a milestone in the overall project.

Activities, and include consortium management and assessment of progress and results. (Please note that your overall approach to management will be described later, in section 2).

Please present your plans as follows<sup>20</sup>:

- i) Describe the overall strategy of the work plan (*Maximum length – one page*)
- ii) Show the timing of the different WPs and their components (Gantt chart or similar)
- iii) Provide a detailed work description broken down into work packages:
  - Work package list (please use table 1.4a);
  - Deliverables list (please use table 1.4b);
  - List of milestones (please use table 1.4c)
  - Description of each work package, and summary (please use table 1.4d);
  - Summary effort table (please use table 1.4e);
- iv) Provide a graphical presentation of the components showing their interdependencies (Pert diagram or similar)
- v) Describe any significant risks, and associated contingency plans

### 1.5 Quality and effectiveness of the S/T Methodology (relevant to CP part only)

Describe the methodology to achieve the objectives of the CP part of the project, especially the way joint PCP will be implemented. This shall include information on the number of phases in the PCP process that are foreseen (this may depend on the complexity of the R&D to be performed), the expected duration/cost of each phase, the number of companies foreseen to be invited to participate in the PCP to have a good representation of possible competing solution paths, etc.

The description of the methodology shall confirm the consortium commitment to:

- use functional specifications for describing the object of the PCP tender (based on jointly specified functional / performance requirements for the solution)
- ensure EU wide publication of the PCP call for tender, at least in English, and evaluate all offers according to the same objective criteria regardless of the geographic location of company head offices, company size or governance structure
- organise the PCP process so as to stimulate participating companies to locate a relevant portion of the R&D and operational activities related to the PCP contract in the European Economic Area or a country having concluded a Stabilisation and Association Agreement with the EU
- use best value for money (and not just lowest price) criteria for evaluating offers
- require participating companies to foresee in their offer a financial compensation according to market conditions compared to exclusive development price for assigning IPR ownership rights to participating companies, in order for the PCP call for tender not to involve State aid
- implement the PCP contracts that will be concluded with the selected organisations in the form of one single framework contract covering all the PCP phases (specific contracts per phase for companies proceeding progressively across the PCP phases)

#### Joint Research Activities (the CP part of project) and associated work plan

A detailed work plan should be presented, broken down into work packages (WPs) which should follow the logical phases of the implementation of the project's Joint Research Activities, and include assessment of progress and results.

Please present your plans as follows:

- i) Describe the overall strategy of the work plan (*Maximum length – one page*)
- ii) Show the timing of the different WPs and their components (Gantt chart or similar)
- iii) Provide a detailed work description broken down into work packages:
  - Work package list (please use table 1.5a);

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<sup>20</sup> The first WP under this section should address the management related activities of the project and the costs relevant to this WP should be reported in the "Management" column of the appropriate A-form. The remaining costs should be reported under the "Co-ordination" column of the appropriate A-form.

- Deliverables list (please use table 1.5b);
  - List of milestones (please use table 1.5c)
  - Description of each work package, and summary (please use table 1.5d);
  - Summary effort table (please use table 1.5e);
  - Cost for the joint pre-commercial procurement per partner (please use table 1.5f)
- iv) Provide a graphical presentation of the components showing their interdependencies (Pert diagram or similar)
- v) Describe any significant risks, and associated contingency plans

Notes:

The number of work packages used must be appropriate to the complexity of the work and the overall value of the proposed project. The planning should be sufficiently detailed to justify the proposed effort and allow progress monitoring by the Commission.

*(Indicative maximum length for the whole of Section 1 – forty pages. This limit does not include the Gantt chart, Pert diagram and tables 1.4 a-e and 1.5 a-f)*

Table 1.4a / 1.5a: *Template - Work package list*

**Work package list**

Work package No <sup>21</sup>	Work package title	Type of activity <sup>22</sup>	Lead partic no. <sup>23</sup>	Lead partic. short name	Person-months <sup>24</sup>	Start month <sup>25</sup>	End month
	<b>TOTAL</b>						

<sup>21</sup> Workpackage number: WP 1 – WP n

<sup>22</sup> Please indicate one activity per work package:

RTD = Research and technological development; COORD = Co-ordination;

MGT = Management of the consortium; SVC = Service activities

<sup>23</sup> Number of the participant leading the work in this work package

<sup>24</sup> The total number of person-months allocated to each work package

<sup>25</sup> Measured in months from the project start date (month 1)

Table 1.4b / 1.5b: Template - Deliverables List

### List of Deliverables

Del. no. <sup>26</sup>	Deliverable name	WP no.	Nature <sup>27</sup>	Dissemination level <sup>28</sup>	Delivery date <sup>29</sup> (proj. month)

<sup>26</sup> 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.

<sup>27</sup> Please indicate the nature of the deliverable using one of the following codes:

**R** = Report, **P** = Prototype, **D** = Demonstrator, **O** = Other

<sup>28</sup> Please indicate the dissemination level using one of the following codes:

**PU** = Public

**PP** = Restricted to other programme participants (including the Commission Services)

**RE** = Restricted to a group specified by the consortium (including the Commission Services)

**CO** = Confidential, only for members of the consortium (including the Commission Services)

<sup>29</sup> Measured in months from the project start date (month 1)

*Table 1.4c / 1.5c: Template - List of milestones*

### List of Milestones

Milestones are control points where decisions are needed with regard to the next stage of the project. For example, a milestone may occur when a major result has been achieved, if its successful attainment is a required for the next phase of work. Another example would be a point when the consortium must decide which of several technologies to adopt for further development.

<b>Milestone number</b>	<b>Milestone name</b>	<b>Work package(s) involved</b>	<b>Expected date<sup>30</sup></b>	<b>Means of verification<sup>31</sup></b>

<sup>30</sup> Measured in months from the project start date (month 1)

<sup>31</sup> Show how you will confirm that the milestone has been attained. Refer to indicators if appropriate. For example: a laboratory prototype completed and running flawlessly; software released and validated by a user group; field survey complete and data quality validated.

Table 1.4d / 1.5d: *Template - Work package description*

**Work package description**

<b>Work package number</b>		<b>Start date or starting event:</b>					
<b>Work package title</b>							
<b>Activity type<sup>32</sup></b>							
<b>Participant number</b>							
<b>Participant short name</b>							
<b>Person-months per participant</b>							

**Objectives**

**Description of work** (possibly broken down into tasks) and role of partners

**Deliverables** (brief description) and month of delivery

<sup>32</sup> Please indicate one activity per work package:

RTD: Research and technological development; COORD: Co-ordination; MGT: Management of the consortium;  
SVC: Service activities



*Table 1.4e / 1.5e: Summary of effort*

**Summary of effort**

A summary of the effort is useful for the evaluators. Please indicate in the table number of person months over the whole duration of the planned work, for each work package by each participant. Identify the work-package leader for each WP by showing the relevant person-month figure **in bold**.

<b>Partic. no.</b>	<b>Partic. short name</b>	<b>WP1</b>	<b>WP2</b>	<b>WP3</b>	<b>...</b>	<b>Total person months</b>
<b>1</b>						
<b>2</b>						
<b>3</b>						
<b>etc</b>						
<b>Total</b>						

*Table 1.5 f: Template - Joint Pre-Commercial Procurement cost table*

Please identify the cost for the joint pre-commercial procurement per partner.

<b>Part. number</b>	<b>Part. short name</b>	<b>Cost (€)</b>
<b>1</b>		
<b>2</b>		
<b>3</b>		
...		
-	<b>Total</b>	

Note that the costs related to the joint pre-commercial procurement implemented under the CP part of the project are considered as a sub-contracting of joint research activities and thus have to be declared under the column “RTD activities” in the relevant A3 forms. The funding of costs for the joint research activities is limited to a maximum of 75% of the eligible costs for the development of the new ICT solutions procured through the joint PCP call for tender (for financing the R&D to be performed by selected tenderers participating in the PCP).

There may be two types of partners contributing to the costs for the joint PCP procurement in Table 1.5.f:

- Type 1 beneficiaries: non-profit public bodies (e.g. for this action public purchasers that are subject to public procurement directive 2004/18/EC that are non-profit public bodies), secondary and higher education establishments or research organisations and SMEs which are unable to identify with certainty their real indirect costs for the project.
- Type 2 beneficiaries: other type of beneficiaries (e.g. for this action this includes private non-SME type public purchasers such as utilities that are subject to public procurement directive 2004/17/EC)

The maximum reimbursement rate for type 1 beneficiaries is 75%. The maximum reimbursement rate for type 2 is 50%. If all partners listed in Table 1.5 f that are contributing financially to the cost of the joint PCP are type 1 beneficiaries, the maximum reimbursement rate for the costs under the CP part is 75%. If all partners listed in Table 1.5 f that are contributing financially to the cost of the joint PCP are type 2 beneficiaries, the maximum reimbursement rate for the costs under the CP part is 50%. If there is a mix of type 1 and type 2 beneficiaries listed in Table 1.5 f that are contributing financially to the cost of the joint PCP, the maximum reimbursement rate for the PCP procurement cost under the CP part will lie in between 50% and 75% and will be calculated taking into account that the maximum reimbursement rates for the financial contribution of type 1 versus type 2 beneficiaries towards the total PCP procurement budget is 75% versus 50% respectively.

## Section 2. Implementation

### 2.1 Appropriateness of the management structure and procedures

Describe the organisational structure and decision-making mechanisms of the project. Show how they are matched to the complexity and scale of the project. This relates in particular to the management structure and procedures proposed for the overall CP-CSA project and for the joint PCP that will be launched during the CP-CSA project.

*(Maximum length for Section 2.1 – five pages)*

### 2.2 Quality and relevant experience of the individual participants

For each participant in the proposed project, provide a brief description of the legal entity, the main tasks they have been attributed, and the previous experience relevant to those tasks. Provide also a short profile of the individuals who will be undertaking the work. This relates in particular to the direct beneficiaries of the EU grant, not to the tenderers that will participate in the PCP (as those are not direct beneficiaries of the EU grant, and they are still unknown at the time of grant signature before the launch of the PCP).

*(Maximum length for Section 2.2 – one page per participant. However, where two or more departments within an organisation have quite distinct roles within the proposal, one page per department is acceptable.*

*The maximum length applying to a legal entity composed of several members, each of which is a separate legal entity (for example an EEIG), is one page per member, provided that the members have quite distinct roles within the proposal.)*

### 2.3 Quality of the consortium as a whole

Describe how the participants collectively constitute a consortium capable of achieving the project objectives, and how they are suited and are committed to the tasks assigned to them. Show the complementarity between participants. Explain how the composition of the consortium is well-balanced in relation to the objectives of the project.

In conformity with the work programme, the composition of the consortium also refers to an appropriate level of representation (local / regional / national) as well as critical mass of public purchasers necessary to trigger wide implementation of the public service innovation strategies and solutions specified and/or developed during the PCP.

**i) Sub-contracting:** If any part of the work is to be sub-contracted by the participant responsible for it, describe the work involved and explain why a sub-contract approach has been chosen for it.

For a good understanding of the conditions on sub-contracting, please read carefully the subcontracting box in section A/3 'budget' of the table in Annex 3 of this guide for applicants.

In conformity with FP7 rules for participation and the work programme, for the CP-CSAs on PCP, the eligible costs for the development of ICT solutions that procured by the beneficiaries from subcontractors selected through the joint PCP call for tender, can be reimbursed up to maximum 75%. This has to be specified carefully in Annex I to the Grant Agreement.

Consortia may identify certain tasks to be subcontracted to external entities or undertaken by in-house consultants under the responsibility of the consortia participants. Such tasks could rely on the services of, for example:

- experts for the PCP tender evaluation committee
- legal experts for assisting in the procedural aspects of the tender

**ii) Other countries:** If a one or more of the participants requesting EU funding is based outside of the EU Member states, Associated countries and the list of International Cooperation Partner Countries<sup>33</sup>, explain in terms of the project's objectives why such funding would be essential.

**iii) Additional partners:** If there are as-yet-unidentified participants in the project, the expected competences, the role of the potential participants and their integration into the running project should be described. (These as-yet-unidentified participants will not be counted in the minimum number of participants for the eligibility of the proposal).

*(No maximum length for Section 2.3 – depends on the size and complexity of the consortium)*

#### **2.4 Appropriateness of the allocation and justification of the resources to be committed**

Describe how the totality of the necessary resources (staff and equipment) will be mobilised, including any resources that will complement the EU contribution. Show how the resources will be integrated in a coherent way, and show how the overall financial plan for the project is adequate. In conformity with the work programme, consortia shall demonstrate that they contain a critical mass of public purchasers with clear financial commitments to undertake a joint PCP.

In addition to the costs indicated on form A3 of the proposal, and the effort shown in section 1.3 above, please identify any other major costs (e.g. equipment). Ensure that the figures stated in Part B are consistent with these.

*(Maximum length for Section 2.4 – two pages)*

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<sup>33</sup> See CORDIS web-site, and annex 1 of the work programme.

## Section 3. Impact

### 3.1 Contribution to the expected impacts listed in the work programme

Describe how your project will contribute towards the expected impacts listed in the work programme in relation to the topic or topics in question (including those impacts listed in objective 11.1 of the work programme). In particular, describe also the expected impacts related to developing a more forward-looking, concerted, public sector approach to societal challenges across boundaries, increasing opportunities for wide market uptake and economies of scale for the supply side and reducing fragmentation of public sector demand for new ICT solutions. Mention the steps that will be needed to bring about these impacts.

Explain why this contribution requires a European (rather than a national or local) approach. Indicate how account is taken of other national or international research activities. Mention any assumptions and external factors that may determine whether the impacts will be achieved.

### 3.2 Dissemination and/or exploitation of project results, and management of intellectual property

Describe the measures you propose for the dissemination and/or exploitation of project results, and how these will increase the impact of the project. In designing these measures, you should take into account a variety of communication means and target groups as appropriate (e.g. policy-makers, interest groups, media and the public at large).

For more information on communication guidance, see [http://ec.europa.eu/research/science-society/science-communication/index\\_en.htm](http://ec.europa.eu/research/science-society/science-communication/index_en.htm)

Describe also your plans for the management of knowledge (intellectual property) acquired in the course of the project.

This relates in particular also to wide publication of results of cross border PCP activities, removal of barriers to market introduction for the developed PCP solutions through joint regulatory action or contribution to standardisation based on jointly defined public sector PCP solution requirements specifications, appropriate consultation of stakeholders and division of IPR rights between public purchasers and companies participating in the PCP according to Appendix 6 of the work programme.

Describe the industrial/commercial involvement to ensure wide exploitation of the results. Describe how the opportunity of involving SMEs has been considered. Pre-commercial procurement is a public procurement of R&D services with the objective to develop breakthrough solutions for public sector problems for which there are no solutions – so typically no large established top tier solution providers and no customer references – on the market yet. It is therefore important to remove unnecessary barriers for innovative new SMEs to make offers. Stringent qualification requirements (e.g. prior customer references) and financial guarantees (e.g. minimum turnover) - which are often difficult for SMEs to comply with - are therefore typically not used as selection criteria in pre-commercial procurements.

*(Maximum length for the whole of Section 3 – ten pages)*

#### Section 4. Ethical Issues

Describe any ethical issues that may arise in their proposal. In particular, you should explain the benefit and burden of their experiments and the effects it may have on the research subject. Identify the countries where research will be undertaken and which ethical committees and regulatory organisations will need to be approached during the life of the project.

The following special issues should be taken into account:

**Informed consent:** When describing issues relating to informed consent, it will be necessary to illustrate an appropriate level of ethical sensitivity, and consider issues of insurance, incidental findings and the consequences of leaving the study.

**Data protection issues:** Avoid the unnecessary collection and use of personal data. Identify the source of the data, describing whether it is collected as part of the research or is previously collected data being used. Consider issues of informed consent for any data being used. Describe how personal identify of the data is protected.

**Use of animals:** Where animals are used in research the application of the 3Rs (Replace, Reduce, Refine) must be convincingly addressed. Numbers of animals should be specified. State what happens to the animals after the research experiments.

**Human embryonic stem cells:** Research proposals that will involve human embryonic stem cells (hESC) will have to address all the following specific points:

- the necessity to use hESC in order to achieve the scientific objectives set forth in the proposal.
- whether the applicants have taken into account the legislation, regulations, ethical rules and/or codes of conduct in place in the country(ies) where the research using hESC is to take place, including the procedures for obtaining informed consent;
- the source of the hESC
- the measures taken to protect personal data, including genetic data, and privacy;
- the nature of financial inducements, if any.

Include the Ethical issues table below. If you indicate YES to any issue, please identify the pages in the proposal where this ethical issue is described. If you are sure that none of the issues apply to your proposal, simply tick the YES box in the last row.

*(No maximum length for Section 4 – depends on the number and complexity of the ethical issues involved)*

#### Notes:

1. For further information on ethical issues relevant to ICT, see Annex 5 of this Guide
2. Only in exceptional cases will additional information be sought for clarification, which means that any ethical review will be performed solely on the basis of the information available in your proposal.
3. A dedicated website that aims to provide clear, helpful information on ethics issues is now available at: [http://cordis.europa.eu/fp7/ethics\\_en.html](http://cordis.europa.eu/fp7/ethics_en.html). The site includes guidance documents on privacy and data protection, developing countries, informed consent procedures etc.

**ETHICAL ISSUES TABLE**

	YES	Page Number
<b>Informed Consent</b>		
• Does the proposal involve children?		
• Does the proposal involve patients?		
• Does the proposal involve persons not able to give consent?		
• Does the proposal involve adult healthy volunteers?		
<b>Biological research</b>		
• Does the proposal involve human genetic material?		
• Does the proposal involve human biological samples?		
• Does the proposal involve human biological data collection?		
• Does the proposal involve human embryos?		
• Does the proposal involve human foetal tissue or cells?		
• Does the proposal involve human embryonic stem cells?		
<b>Privacy</b>		
• Does the proposal involve processing of genetic information or personal data (e.g. health, sexual lifestyle, ethnicity, political opinion, religious or philosophical conviction)		
• Does the proposal involve tracking the location or observation of people without their knowledge?		
<b>Research on Animals</b>		
• Does the proposal involve research on animals?		
• Are those animals transgenic small laboratory animals?		
• Are those animals transgenic farm animals?		
• Are those animals cloned farm animals?		
• Are those animals non-human primates?		
<b>Research Involving Third Countries</b>		
• Is any part of the research carried out in countries outside of the European Union and FP7 Associated states?		
<b>Dual Use</b>		
• Does the research have direct military application		
• Does the research have the potential for terrorist abuse		
<b>ICT Implants</b>		
• Does the proposal involve clinical trials of ICT implants?		
<b>(IF NONE) I CONFIRM THAT NONE OF THE ABOVE ISSUES APPLY TO MY PROPOSAL</b>		

## ***Annex 5: Ethical Guidelines for undertaking ICT research in FP7***

### **1. Introduction**

In recent years there has been an increase in the importance of ethical issues related to ICT research and technological developments.

The decision of the European Parliament and the Council concerning FP7<sup>34</sup> states that research activities supported by the Framework Programme should respect fundamental ethical principles, including those reflected in the Charter of Fundamental Rights of the European Union<sup>35</sup> and take into account opinions of the European Group on Ethics in Science and New Technologies (EGE)<sup>36</sup>.

Article 15 of the FP7 draft rules of participation<sup>37</sup> states that any proposal which contravenes fundamental ethical principles or which does not fulfil the conditions set out in the specific programme, the workprogramme or in the call for proposals shall not be selected and may be excluded from the evaluation, selection and award procedures at any time.

Applications for EU-funded research activities may, if appropriate, include specific tasks or a specific work package that explicitly addresses ethical concerns (in terms of the research, its conduct and outcomes) and outlines how ethical issues raised by the proposed research will be handled.

The purpose of this guidance is to assist proposers in identifying potential ethical issues arising from the proposed ICT research.

### **2. Conduct of ICT Research**

All research areas within ICT of FP7 may raise ethical issues of varying seriousness. Some proposals will be more sensitive than others. It is likely that new, sensitive applications will come to the fore during the term of FP7.

#### **2.1 A responsible approach**

It is likely that most of the principles of the Charter of Fundamental Rights of the European Union<sup>38</sup> will be relevant to the approach adopted by ICT researchers. These principles cover dignity, freedom, equality, solidarity, citizens' rights and justice. Proposals must comply with Article 8 of the European Human Rights Convention<sup>39</sup>. In particular, given the pervasive and ubiquitous nature of ICT and the many opportunities it offers, researchers should consider the sensitive implications of their proposals for privacy and autonomy.<sup>40</sup> However, researchers should recognise that new dangers associated with the process of ICT research can exist. They should carry out a prior assessment of risk and identification of precautionary actions proportional to the potential risk/harm.<sup>41</sup>

Researchers have a duty to alert public authorities to the ethical and practical implications of the ICT research outcomes, as and when particular issues become apparent within the research process.<sup>7</sup>

Researchers should comply with national legislation, European Union legislation, respect international conventions and declarations and take into account the Opinions of the European Group on Ethics. However, consideration of ethical issues goes beyond simple compliance with current regulations and laws.

#### **2.2 Privacy and informed consent**

The right to privacy and data protection is a fundamental right<sup>42</sup> and therefore applicable to ICT research.

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<sup>34</sup> Decision 1982/2006/EC: Official Journal L412 of 18/12/06

<sup>35</sup> [http://www.europarl.europa.eu/charter/default\\_en.htm](http://www.europarl.europa.eu/charter/default_en.htm)

<sup>36</sup> The EGE is an independent, multidisciplinary body, appointed by the Commission to examine ethical questions arising from science and new technologies and on this basis to issue *Opinions* - [http://ec.europa.eu/european\\_group\\_ethics/index\\_en.htm](http://ec.europa.eu/european_group_ethics/index_en.htm)

<sup>37</sup> Official Journal L391 of 30/12/06

<sup>38</sup> The Charter of Fundamental Rights of the European Union - [http://www.europarl.europa.eu/charter/pdf/text\\_en.pdf](http://www.europarl.europa.eu/charter/pdf/text_en.pdf)

<sup>39</sup> <http://conventions.coe.int/treaty/en/Treaties/Html/005.htm>

<sup>40</sup> Opinion 10 of EGE - The Ethical Aspects of the 5<sup>th</sup> Framework Programme , [http://ec.europa.eu/european\\_group\\_ethics/docs/opinion10\\_en.pdf](http://ec.europa.eu/european_group_ethics/docs/opinion10_en.pdf)

<sup>41</sup> Opinion 20 of EGE – Ethical Aspects of ICT Implants in the Human Body - [http://ec.europa.eu/european\\_group\\_ethics/docs/avis20\\_en.pdf](http://ec.europa.eu/european_group_ethics/docs/avis20_en.pdf)

<sup>42</sup> The Charter of Fundamental Rights of the European Union - [http://www.europarl.europa.eu/charter/pdf/text\\_en.pdf](http://www.europarl.europa.eu/charter/pdf/text_en.pdf)



Researchers must be aware that volunteers<sup>43</sup> have the right to remain anonymous<sup>44</sup>. Researchers must comply with Data Protection legislation<sup>45</sup> in the Member State where the research will be carried out regarding ICT research data that relates to volunteers.

Informed consent is required whenever ICT research involves volunteers in interviews, behavioural observation, invasive and non-invasive experimentation, and accessing personal data records. The purpose of informed consent is to empower the individual to make a voluntary informed decision about whether or not to participate in the research based on knowledge of the purpose, procedures and outcomes of the research.

Before consent is sought, information must be given specifying the alternatives, risks, and benefits for those involved in a way they understand. When such information has been given, free and informed consent must be obtained. Depending on the nature of the research, different consent procedures may be used. Special consideration must be given when volunteers have reduced autonomy or are vulnerable<sup>3</sup>.

The majority of European citizens view personal privacy as an important issue. Research, for example, on RFID<sup>46</sup> and ICT for healthcare<sup>47</sup>, is likely to raise privacy issues. Therefore, researchers must ensure that the manner in which research outcomes are reported does not contravene the right to privacy and data protection. Furthermore, researchers must carefully evaluate and report the personal privacy implications of the intended use or potential use of the research outcomes. Wherever possible, they must ensure that research outcomes do not contravene these fundamental rights.

### 2.3 Use of animals in ICT research

In accordance with the Amsterdam protocol on animal protection and welfare, animal experiments must be replaced with alternatives wherever possible. Suffering by animals must be avoided or kept to a minimum. This particularly applies to animal experiments involving species which are closest to human beings<sup>48</sup>. Thus ICT research involving animals should conform to the ethical principles of replacement, reduction, refinement and minimisation of suffering<sup>3</sup>.

Proposers must carefully justify animal experiments in cross-science proposals for non-medical objectives. Furthermore, they should identify the scientific areas which would benefit from knowledge gained through animal experiments. Proposers must be aware that Member States may have differing and possibly conflicting interpretations of animal welfare in research, and the research must meet regulations in the country in which it will be carried out.

## 3 Specific guidance in some currently sensitive areas

### 3.1 ICT implants<sup>49</sup> and wearable computing

- ICT implants should only be developed if the objective cannot be achieved by less-invasive methods such as wearable computing devices and RFID tags.
- To the extent that an individual, via an ICT implant or wearable computing device, becomes part of an ICT network, the operation of this whole network will need to respect privacy and data protection requirements.
- ICT implants in healthcare are, in general, acceptable when the objective is saving lives, restoring health, or improving the quality of life. They should be treated in the same way as drugs and medical devices.<sup>50</sup>
- ICT implants to enhance human capabilities should only be developed: to bring individuals into the “normal” range for the population, if they so wish and give their informed consent; or to improve health prospects such as enhancing the immune system. Their use should be based on need, rather than economic resources or social position.

<sup>43</sup> “Volunteers” is used to describe all those who are the subjects of research observations, experiments, tests etc.

<sup>44</sup> Opinion 10 of EGE - The Ethical Aspects of the 5<sup>th</sup> Framework Programme ,  
[http://ec.europa.eu/european\\_group\\_ethics/docs/opinion10\\_en.pdf](http://ec.europa.eu/european_group_ethics/docs/opinion10_en.pdf)

<sup>45</sup> National legislation transposing Directive 95/46/EC -  
[http://ec.europa.eu/justice\\_home/fsj/privacy/docs/95-46-ce/dir1995-46\\_part1\\_en.pdf](http://ec.europa.eu/justice_home/fsj/privacy/docs/95-46-ce/dir1995-46_part1_en.pdf)

<sup>46</sup> RFID Technology - Results of the Public Consultation on Article 29 Working Document 105 on Data Protection Issues Related to RFID Technology Adopted on 28 September 2005  
[http://europa.eu.int/comm/justice\\_home/fsj/privacy/workinggroup/consultations/rfid\\_en.htm](http://europa.eu.int/comm/justice_home/fsj/privacy/workinggroup/consultations/rfid_en.htm)

<sup>47</sup> Opinion 13 of EGE - Ethical Issues of Healthcare in The Information Society.-  
[http://ec.europa.eu/european\\_group\\_ethics/docs/avis13\\_en.pdf](http://ec.europa.eu/european_group_ethics/docs/avis13_en.pdf)

<sup>48</sup> Council Directive on Protection of Animals used for Experimental and other Scientific Purposes  
[http://europa.eu.int/comm/food/fs/aw\\_legislation/scientific/86-609-eeec\\_en.pdf](http://europa.eu.int/comm/food/fs/aw_legislation/scientific/86-609-eeec_en.pdf)

<sup>49</sup> Opinion 20 of EGE – Ethical Aspects of ICT Implants in the Human Body -  
[http://ec.europa.eu/european\\_group\\_ethics/docs/avis20\\_en.pdf](http://ec.europa.eu/european_group_ethics/docs/avis20_en.pdf)

<sup>50</sup> Such research is partly covered by Council Directive 90/385/EEC relating to active implantable medical devices-  
[http://europa.eu.int/eur-lex/en/consleg/pdf/1990/en\\_1990L0385\\_do\\_001.pdf](http://europa.eu.int/eur-lex/en/consleg/pdf/1990/en_1990L0385_do_001.pdf)

- ICT implants or wearable computing devices must not: allow individuals to be located on a permanent and/or occasional basis, without the individual's prior knowledge and consent; allow information to be changed remotely without the individual's prior knowledge and consent; be used to support any kind of discrimination; be used to manipulate mental functions or change personal identity, memory, self-perception, perception of others; be used to enhance capabilities in order to dominate others, or enable remote control over the will of other people.
- ICT implants should not be developed to influence future generations, either biologically or culturally.
- ICT implants should be developed to be removed easily.

### 3.2 eHealth<sup>51</sup> and genetics

Personal health data must be treated as 'sensitive personal data'<sup>52</sup>. ICT researchers using it have a duty of confidentiality equivalent to the professional duty of medical secrecy. Therefore:

- The use of personal health data in ICT research for the purposes from which society as a whole benefits must be justified in the context of the personal rights.
- The security of ICT in healthcare is an ethical imperative to ensure the respect for human rights and freedoms of the individual, in particular the confidentiality of data and the reliability of ICT systems used in medical care.
- Proposers should be particularly aware when ICT is linked to sensitive medical areas such as the use of genetic material<sup>1</sup>.
- Proposers should access established general medical and genetics ethical guidance when formulating their proposals.

### 3.3 ICT and Bio/Nano-electronics

ICT-bio/nano-electronics has a strong potential for mis-use. Consequently, proposers should pay particular attention to the guidelines in Section 2 in this area<sup>53</sup>.

- Researchers involved in ICT-bio/nano-electronics research proposals should be aware that certain applications, e.g. miniaturised sensors, may have specific implications for the protection of privacy and personal data<sup>4</sup>.
- ICT-bio/nano-electronics research may overlap with other scientific disciplines such as biology. In these situations proposers should draw upon the ethical guidance of that discipline.

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<sup>51</sup> Opinion 13 of EGE - Ethical Issues of Healthcare in The Information Society. - [http://ec.europa.eu/european\\_group\\_ethics/docs/avis13\\_en.pdf](http://ec.europa.eu/european_group_ethics/docs/avis13_en.pdf)

<sup>52</sup> Directive 95/46/EC -

[http://ec.europa.eu/justice\\_home/fsj/privacy/docs/95-46-ce/dir1995-46\\_part1\\_en.pdf](http://ec.europa.eu/justice_home/fsj/privacy/docs/95-46-ce/dir1995-46_part1_en.pdf)

<sup>53</sup> COM (2004) 338 final - [http://ec.europa.eu/prelex/rech\\_simple.cfm?CL=en](http://ec.europa.eu/prelex/rech_simple.cfm?CL=en)



Proposal summary

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The Commission services will reply by electronic mail giving a brief assessment of this pre-proposal. The assessment does not constitute in any respect a pre-evaluation of the proposal in terms of scientific and technical quality. The advice given by the Commission is strictly informal and non-binding. The advice provided does not in any way engage the Commission with regard to acceptance or rejection of the proposal when it is finally submitted.