These information sheet and informed consent templates were provided by CIREP, but certain fields have been modified to reflect the approved BESLab protocol. The text proposed is tentative and can be modified depending on the project’s characteristics. Footnotes are informative: please take them into consideration and eliminate them before saving the final version of the document.

**Information Sheet**

**Title of the research project:** [Project title]

**Principal Investigator:** Prof./Dr. XYZ, [work address], xyz@upf.edu

**Institution:** Universitat Pompeu Fabra [or name of the relevant institution]

**Funding agency:** This project is funded by [funding body, award/grant id].

**Objectives and duration of the project:** The goal of this study is [please describe the goal of your project without revealing details or hypotheses that may bias participants’ responses]. The duration of this project is [please indicate the duration of the project].

**Methodology and participation:** [Describe the project’s methodology, especially the procedures related to subjects’ participation. Indicate clearly the estimated time to complete the participation, the number of sessions, the tasks to be carried out and where they will take place, as well as any other important details.] The experiment will take place in the BESLab premises, located in room 24.320 of the Mercè Rodoreda building.

**Privacy:** [Indicate the measures adopted to preserve participants’ privacy, e.g. *In order to protect your privacy, we will not identify your data with your name, but rather with a code that will only be known to the research team members. In order to make data only accessible to research team members, physical data will be stored in a locked secure location and digital data will be stored with access control systems. In the event of data publication, only anonymous data will be published. Anonymized data may be hosted or published in a public repository.*]

[Please choose option A, B, or C as relevant and delete the other options]

**[A.]** We need to process your personal data for payment purposes. To that end, your data will be shared with UPF administrative staff and will be treated separatedly and independently from your experiment responses. The data collected during the experiment cannot be used to identify you. Thus, once the experiment concludes, we will not be able to delete the data you generated, should you ask for it, unless you can provide us with additional information that allows us to identify your contribution.

Only anonymized data will be published. Anonymized data may be reused for other research projects or archived or published in a public repository.

Payment related data will be kept for the duration of the project and for its scientific validation and according to UPF regulations.

**[B.]** We need to process your identifying personal data for payment purposes. These data allow us to identify the responses you provide during the experiment. However, the payment related data will be shared with UPF administrative staff and will be treated separatedly and independently from your experiment responses.

[Please add option 1 or 2 as needed and complete if necessary.]

**[Option 1]** Your experiment responses will be stored on a protected UPF server and will only be accessible to the project team members and the lab manager.

**[Option 2]** Your experiment responses will be stored on [software name] servers, which are GDPR compliant and, later, on a protected UPF server. They will only be accessible to the project team members and the lab manager.

Only anonymized data will be published. Anonymized data may be reused for other research projects or archived or published in a public repository.

Informed consent sheets will be kept in a lock cabinet and will only be accessible to the principal investigator.[[1]](#footnote-2)

Personal data will be kept for the duration of the project and its scientific validation and according to UPF regulations.

**[C.]** We need to process personal data that might identify you. Therefore, they will be treated in accordance with personal data protection regulations.

[Please add option 1 or 2 as needed and complete if necessary.]

**[Option 1]** Your experiment responses will be stored on a protected UPF server and will only be accessible to the project team members and the lab manager.

**[Option 2]** Your experiment responses will be stored on [software name] servers, which are GDPR compliant and, later, on a protected UPF server. They will only be accessible to the project team members and the lab manager.

We will also need to process your personal data for payment purposes. The data will be shared with UPF administrative staff and will be treated separatedly and independently from your experiment responses.

Only anonymized data will be published. Anonymized data may be reused for other research projects or archived or published in a public repository.

Informed consent sheets will be kept in a lock cabinet and will only be accessible to the principal investigator.[[2]](#footnote-3)

Personal data will be kept for the duration of the project and its scientific validation and according to UPF regulations.[[3]](#footnote-4)

**Compensation:** Your participation will be compensated [indicate how. Specify if compensation requires complete participation.] Partial participation will not be compensated if the participant decides to withdraw from the session. However, if the session is interrupted due to computer malfunctioning, compensation will be prorated.

**Risks and benefits:** [Describe the risks associated to participation. If there are no specific risks, you may use the following text: *Participating in this study does not entail risks greater than those ordinarily encountered in daily life.* Describe benefits associated to participation. If no benefits have been identified, you may use the following text: *We cannot and do not guarantee that you will receive any benefits from this study*.]

**Voluntary nature of participation:** Your participation in this study is on a voluntary basis and you may withdraw from the study at any time without having to justify why.

**Contact information:** If you have any question about this study, you may contact the IP (Prof. XYZ, [xyz@upf.edu](mailto:xyz@upf.edu)) or [contact information of another research team member].

If you have doubts, complaints, or questions about this study or about your rights as a research participant, you may contact UPF’s Institutional Committee for the Ethical Review of Projects (CIREP) by phone (+34 93 542 21 86) or by email ([secretaria.cirep@upf.edu](mailto:secretaria.cirep@upf.edu)). CIREP is not part of the research team and will treat any information you send confidentially.

[If UPF is responsible for the personal data processing, please include this information note at the end of the information sheet:

*In accordance with the General Data Protection Regulation (GDPR) 2016/679 (EU), we provide the following information:*

***Data controller:*** *Universitat Pompeu Fabra. Pl. de la Mercè, 10-12. 08002 Barcelona. Tel. +34 93 542 20 00. You can contact UPF’s Data Protection officer by sending an email to* [*dpd@upf.edu*](mailto:dpd@upf.edu)*.*

***Purposes of the processing:*** *Carrying out the above-mentioned research project. Personal data will be kept during the execution of the project and for two more years after its conclusion for scientific validation.[[4]](#footnote-5)*

***Legal basis:*** *Data owner’s consent. You can withdraw your consent at any time.*

***Recipients (option 1):*** *[[5]](#footnote-6) Your personal data will only be processed by Universitat Pompeu Fabra and will not be transferred to third parties without your consent, except as otherwise provided by law. Data may be anonymized and published in an open science repository.*

***Recipients (option 2):*** *Your personal data will be processed by Universitat Pompeu Fabra and…[[6]](#footnote-7) Data may be anonymized and published in an open science repository.*

***Rights:****You can access your data; request their rectification, deletion, and in certain cases their portability; you may object to their processing and apply for their limitation by following the procedures described at* [*www.upf.edu/web/proteccio-dades/drets*](http://www.upf.edu/web/proteccio-dades/drets)*. You can contact UPF’s Data Protection officer (*[*dpd@upf.edu*](mailto:dpd@upf.edu)*) for any queries or if you feel that your rights are not properly respected. Should you not be satisfied, you may file a complaint with the Catalan Data Protection Authority.*]

**Informed Consent Form**

**Title of the research project:** [Project title]

**Principal investigator:** Prof./Dr. XYZ, [work address], [xyz@upf.edu](mailto:xyz@upf.edu)

**Institution:** Universitat Pompeu Fabra [or the name of the relevant institution]

I HEREBY CONFIRM that:

* I have read the information sheet regarding the research project,
* I have been able to formulate questions on the project,
* I have received enough information on the project,
* I am between […] and […] years old.[[7]](#footnote-8)

I UNDERSTAND that my participation is voluntary and that I can withdraw from or opt out of the study at any time without any need to justify my decision.

☐ I GIVE MY CONSENT to participate in this study.

☐[[8]](#footnote-9) I GIVE MY CONSENT to […].[[9]](#footnote-10)

Name and last name(s):  
Signature:  
Place and date:

[If UPF is responsible for the personal data processing, please include the same information note you included at the end of the information sheet here.]

1. This sentence should only be added if hard copies of the consent form are used. [↑](#footnote-ref-2)
2. This sentence should only be added if hard copies of the consent form are used. [↑](#footnote-ref-3)
3. Modify this statement as necessary for your project needs. [↑](#footnote-ref-4)
4. The retention period can be adapted to the project’s needs. [↑](#footnote-ref-5)
5. Choose the most convenient option (1 or 2). [↑](#footnote-ref-6)
6. If necessary, add other recipients (specific ones or categories). Contracts with specific clauses on data protection will have to be signed. Types of recipients:

   companies hired to provide a service which requires personal data processing (for example, fieldwork);

   collaborating entities that have access to or process personal data (for example, collaborative projects involving several universities);

   third parties that could have access to the data: for example, in case of image publications for a presentation, scientific publications, or the project’s web page, or in case of possible secondary use of personal data for future projects (explicit consent is needed).

   It is mandatory to indicate data transfers outside the European Economic Area (where the General Data Protection Regulation and its requirements apply) and the requirements established by the Regulation must be complied with. In addition, publication of data online is considered a type of data transfer and participants need to be duly informed about it. [↑](#footnote-ref-7)
7. Please indicate the age range according to the project’s inclusion criteria. [↑](#footnote-ref-8)
8. Add checkboxes so that participants can select the options they want when all of them are not required to carry out the research project. [↑](#footnote-ref-9)
9. Add all the types of processing that require explicit consent from the participant. Options include:

   processing of special categories of data (indicate the types of data that will be processed),

   reuse of my personal data for other projects by the same UPF research group,

   transmission of my personal data to be used in projects in the same field carried out by different research groups,

   transfer of my personal data to countries outside the European Economic Area,

   publication of identifying data (such as my image or voice). [↑](#footnote-ref-10)