**The Informed Voluntary Consent Form (IVCF) should include at least the following headings in a way that the volunteer can understand (expressed in Turkish, avoiding medical terms as much as possible):**

1. That the study is a research,
2. Purpose of the research,
3. All of the methods to be followed or applied to the volunteer, including the methods to be applied during the research process,
4. Responsibilities of the volunteer,
5. Experimental parts of the study, if any (informed consents specific to the experimental and control groups),
6. Risks or discomfort to which the volunteer (embryo, fetus or infant if the research will be conducted on pregnant or postpartum women) will be exposed,
7. When there is no targeted benefit for the volunteer regarding the expected benefits of the research, the volunteer is informed about this situation,
8. If available, information on payments to volunteers for expenses such as transport, meals, etc,
9. The volunteer's participation in the research is voluntary and the volunteer can refuse to participate in the research or withdraw from the research at any time, without being exposed to any penalty or sanction, and without losing any rights,
10. Monitors, examiners, Ethics Committee, Institution and other relevant health authorities may have direct access to the volunteer's original medical records, but this information will be kept confidential, and the volunteer or his/her legal representative will have allowed such access by signing the written informed consent form,
11. In accordance with the relevant legislation, the records that will reveal the identity of the volunteer will be kept confidential and cannot be disclosed to the public; even if the research results are published, the identity of the volunteer will remain confidential,
12. The volunteer or his/her legal representative will be informed in a timely manner when new information is obtained that is relevant to the subject of the research and that may affect the volunteer's willingness to continue to participate in the research,
13. Contact persons for the volunteer to obtain further information about the research and about their rights, and telephone numbers where they can be reached 24 hours a day,
14. Situations or reasons that require the termination of the volunteer's participation in the research,
15. The period foreseen for the volunteer to continue the research,
16. The estimated number of volunteers expected to participate in the survey,
17. *“I have read all the explanations in the Informed Consent Form. Written and verbal explanations regarding the research whose subject and purpose were stated above were made to me by the responsible researcher named below. “I know that I participated in the research voluntarily and that I can withdraw from the research at any time, with or without reason.” containing similar expressions,*
18. “I agree to participate in the research in question with my consent, without any pressure or coercion.” containing similar expressions,
19. The volunteer's name/surname/signature/date must be included,
20. The name/surname/signature/date of a competent researcher in the research team must be included,
21. If necessary, the name/surname/signature/date of the person who witnessed the consent process should be included,
22. If necessary, the name/surname/signature/date of the legal representative should be included,
23. To research biological materials obtained from volunteers; “My biological samples (blood, urine, etc.) taken within the scope of the [Public Name of the Research] research; Information should be included with the appropriate statement marked as "I only allow it to be used in the research mentioned above" or "I allow it to be used in all future research" or "I do not allow it to be used under any circumstances".
24. IVCF cannot contain any provision or statement that would eliminate the legal rights of the volunteer or his legal representative. In addition, it cannot contain any provision or statement that would relieve the researcher and the institution from any liability arising from the sponsor's or their representatives' negligence.