**Waiver or Alternation of Documentation of Informed Consent Explanatory Statement**

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| --- | --- | --- | --- |
| **\* Receipt confirmation** (This is a field for Committee receivers, please researchers keep it blank.) | | | |
| **Receipt No.** |  | **Date of receipt** |  |

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| --- | --- | --- | --- | --- | --- | --- | --- |
| **1. Basic Information** | | | | | | | |
| **Project Name** | | (Korean) | | | | | |
| (English) | | | | | |
| **Type Research** | | □ Human subjects research → method : □ qualitative research □ quantitative research  □ Human materials research □ Embryo research □ Other( ) | | | | | |
| **Principal**  **Investigator** | | Name | (Korean) | | (English) | | |
| Position | □ Professor □ student (□ Master’s course □ Doctors course)  □ Other : Post-Doc, ect. | | | | |
| Affiliation |  | | | | |
| Tel. |  | e-mail | |  | |
| **2.**  **Written consent exemption** | | | | | | | |
| **2-1. Requirement: Only if all entries are under ‘Yes’, written consent exemption is possible.** | | | | | | | |
| 1 | If waive written consent, the risk to human subjects and its derivatives of donors is extremely low. | | | | | | □ Yes □ No |
| 2 | In the case to write consent, it is the only record of human subjects and its derivatives of donor participation. | | | | | | □ Yes □ No |
| 3 | If the consent is exposed with problems of confidentiality, it may cause damage such as stigma, civil and criminal liability. | | | | | | □ Yes □ No |
| 4 | Human subjects and its derivatives of donor don’t want to write consent for these reasons. | | | | | | □ Yes □ No |
| 5 | Although there is no written but verbal agreement is provided with the necessary procedures to receive consent. | | | | | | □ Yes □ No |
| **2-2. Detailed explanation to receive written consent exemption** | | | | | | | |
|  | | | | | | | |
| **2-3. Verbal consent procedure** | | | | | | | |
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* All of the information filled in this form is consistent with the contents of the research to perform
* In the case to confirm that written consent exemption is not possible from Commission, I will apply deliberation with related documents to get written consent. I will start research after receiving approval to use the legal consent..

I submit waiver or alternation of documentation of informed consent explanatory statement as below.

(Date and signature must be written by hand.)

Submission Date: DD/MM/YYYY

Research Director: (Sign)

Academic advisor: (Sign)