**APPENDIX 7**

**OKLAHOMA CHRISTIAN UNIVERSITY**

 **REQUEST FOR WAIVER OF CONSENT**

Instructions: If you are requesting a waiver of informed consent or a waiver of the consent procedure requirement to include all or alter some or all the elements of informed consent [45CFR46.117(c)], you must document the responses to each of the statements, citing supporting sections of the study protocol.

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| **Responsible Project Investigator (RPI)** |
| 1)RPI First Name | RPI Last Name |
|  |
| 2) Project Title |
|  |
| 3) The research in its entirety involves no greater than minimal risk.\_Yes\_No4) The waiver of informed consent will not adversely affect the rights and welfare of the subjects.\_Yes\_No5) It is not practicable to conduct the research without the waiver/alteration.\_Yes\_No6) Whenever appropriate, subjects will be provided with additional pertinent information after their participation.\_Yes\_No |
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| 7) If you have selected the “yes” response to each of the four statements above, in order to receive the waiver, you must:* Describe the reason(s) why the waiver is necessary and explain whether the entire informed consent is being waived or only certain required elements are being waived. (If so, list which ones)

Note: If a waiver is granted under the above conditions, documentation of informed consent (i.e., signed consent form) is also waived. Even if the waiver is granted, the IRB may require other conditions. The IRB may require the researcher to provide subjects with an informed consent sheet (written summary/notification document) about the research. |
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