Hello and thank you for accessing this form from the University of Maryland, Baltimore County Institutional Review Board web site.

Prior to submitting, please ensure that spelling and grammar are correct; this will assist in the timely review of this form during the IRB evaluation process.

Enter information by clicking the ☐ box or typing in the **Click here to enter text.** area.

**Please go to UMBC** [**IRB website**](http://research.umbc.edu/institutional-review-board-human-subjects/) **for all up-to date guidance and information regarding the below questions.**

Protocol Approval Number: Click here to enter text.

Protocol Title:  Click here to enter text.

Investigator(s): Click here to enter text.

In order to completely waive the informed consent process, please provide a response to the following questions as they apply to your full board reviewed research. To qualify for IRB approval, you must provide a response to **ALL** of the following questions. Please be specific in explaining why each statement is applicable to your research.

|  |  |
| --- | --- |
| **1. The research in its entirety involves no more than “minimal risk” to participants.** | [ ]  Yes [ ]  No |
| *Explain:*       |
| **2. The waiver will not adversely affect the rights and welfare of the participants.** | [ ]  Yes [ ]  No |
| *Explain:*       |
| **3. The research could not be practically carried out without the waiver.** | [ ]  Yes [ ]  No |
| *Explain:*       |
| **4. Whenever appropriate, participants will be provided with additional pertinent information after participation.** | [ ]  Yes [ ]  No |
| *Explain:*       |

**If yes, please provide justification for waiving documentation of consent that is specific to your study:** Click here to enter text.

*[Please explain how, in the absence of signed written consent forms, consent will be documented, e.g. tape recordings, videos, chart notes, etc.]* **This justification MUST be included in the “Consent” section of the IRB application.**

**Check the below on which process will be used to record consent:**

[ ]  Audio recording

[ ]  Video recording

[ ]  Web based (on-line) consent acknowledgement box

[ ]  Information Sheet

[ ]  Other(s) - Please specify: Click here to enter text.

**Electronically submit the protocol and any accompanying documents to** [**irbsubmissions@umbc.edu**](file:///%5C%5Csharedvol.ad.umbc.edu%5CDept%5CORA%5CHARPO%5CIRB%5Cirb%20forms%5Cirbsubmissions%40umbc.edu)**.**

By typing your name, email address and date, the investigator(s) certify they will abide by all UMBC IRB policies and procedures and understand that no research activities will be conducted with human participants prior to obtaining the required approvals.

Investigator’s Signature: Click here to enter text. Email: Click here to enter text. Date: Click here to enter a date.

Investigator’s Signature: Click here to enter text. Email: Click here to enter text. Date: Click here to enter a date.

Faculty Advisor's Signature: Click here to enter text. Email: Click here to enter text. Date: Click here to enter a date.

***\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_***

**IRB Action**

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Approved, IRB Chair/Date

(request for waivers of consent form) – 06/13/2017