



# Mohave Community College Institutional Review Board

## Individual Investigator Agreement

*A non-MCC researcher uses this form to certify that they will adhere to all applicable laws, regulations, policies, and IRB determinations. Individual Investigators must sign this statement of commitment to MCC's Human Subject Protection Policies and IRB oversight for each applicable research protocol.*

<b>Institution providing IRB Review</b>		
<b>Mohave Community College</b>		
IRB Registration #: Click or tap here to enter text.		
Federal-wide Assurance (FWA) for the Protection of Human Subjects	Number: Click or tap here to enter text.	

<b>Individual Investigator Relying on Designated IRB</b>		
Name: Click or tap here to enter text.		
Institution: Click or tap here to enter text.		
Project Title: Click or tap here to enter text.		

## Individual Investigator Agreement

The above-named Individual Investigator has reviewed:

### 1. Ethical and Regulatory Guidelines

- The Belmont Report: Ethical Principles and Guidelines for the Protection of Human Subjects of Research*
- The U.S. Department of Health and Human Services (HHS) regulations for the protection of human subjects at *45 CFR part 46*
- The *Federal wide Assurance (FWA)* and applicable Terms of the FWA for the institution referenced above
- The relevant *MCC IRB policies* for the protection of human subjects

### 2. Institutional Agreement:

The Investigator has read and agrees to all applicable terms of the *IRB Authorization and Services Agreement* between their Institution and MCC.

### 3. Compliance Responsibility:

The Investigator understands and hereby accepts the responsibility to comply with the standards, requirements, and directives stipulated in this Agreement and by the *MCC Institutional Review Board*



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(MCC IRB) to protect the rights and welfare of human subjects involved in research conducted under this Agreement.

**4. Legal and Policy Compliance:**

The Investigator will comply with all other applicable federal, international, state, and local laws, regulations, and policies that may provide additional protection for human subjects participating in research conducted under this Agreement.

**5. MCC IRB Authority:**

The Investigator agrees to comply with all MCC IRB determinations, acknowledging its final authority over research approvals.

**6. Required Training:**

The Investigator and their research sponsor (if applicable) will complete any *educational training* required by the MCC IRB prior to initiating research covered under this Agreement.

**7. Protocol Changes:**

The Investigator will report promptly to the MCC IRB any proposed changes in the research conducted under this Agreement. The Investigator will not initiate changes in the research without prior MCC IRB review and approval, except where necessary to eliminate apparent immediate hazards to subjects.

**8. Unanticipated Problems:**

The Investigator will report immediately (within *seven calendar days*) to the MCC IRB any *unanticipated problems* involving risks to subjects or others in research covered under this Agreement.

**9. Informed Consent:**

When responsible for enrolling subjects, the Investigator will obtain, document, and maintain records of *informed consent* for each subject or their legally authorized representative as required by federal regulations and stipulated by the MCC IRB.

**10. Ongoing Review and Reporting:**

The Investigator acknowledges and agrees to cooperate in the MCC IRB's responsibility for *initial and continuing review, record keeping, reporting, and certification* for the research referenced above. The Investigator will provide all information requested by the MCC IRB in a timely manner.

**11. Subject Enrollment:**

The Investigator will not enroll subjects in research under this Agreement prior to its *review and approval* by the MCC IRB.

**12. Emergency Medical Care:**

Emergency medical care may proceed without MCC IRB approval when permitted by federal regulations and state law.




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### 13. Additional Research Participation:

This Agreement does not preclude the Investigator from taking part in research not covered by this Agreement.

### 14. Primary Responsibility for Human Subjects:

The Investigator acknowledges that they are *primarily responsible* for safeguarding the rights and welfare of each research subject and that the subject's rights and welfare must take precedence over the goals and requirements of the research.

Individual investigator
Full address: Click or tap here to enter text.
Phone Number: Click or tap here to enter text.
Email: Click or tap here to enter text.
Signature:   <hr/>