The University of Ver

Institutional Review Board for Human Subject Research

If applicable, are reproductive risks adequately described and is appropriate birth control language included? If pregnant participants or partners are to be followed, include a sepearte consent form	
Anticipated Benefits	
Describe all potential benefits in the consent including knowledge to help others in the future.	
Data Management Plan/Confidentiality	
Ensure your research plan adequately provides for monitoring the data collected, ensuring the confidentiality of the subjects.	

Other considerations:

Have you completed CITI training? This required online program consists of ethics training modules for research with human subjects.

Students – Have you contacted and worked with your faculty advisor and has the protocol been approved by your advisor? Please note, your Faculty Sponsor will be designated within your Click submission, and they must complete their assigned Ancillary Review prior to the IRB's review of your study materials.

Are you requesting reliance on another IRB? Please review our reliance page prior to submission.

Any potential conflicts of interest? Please ensure you have completed a COI disclosure with UVM

Do you plan to share, receive or transfer research materials outside of UVM, such as tissues, cell lines or devices? Don't forget to request a Material Transfer Agreement.

Need an agreement, contract or data use agreement? UVM Sponsored Programs can help. Data use agreements and contracts through the hospital can contact