

Check List for IRB Application Completion
CLINICAL RESEARCH CENTER
MIAMI VALLEY HOSPITAL

- WSU IRB Petition Form: All questions answered with Signatures
- Protocol (This is a written description of the work proposed. It should include sample data collection sheets and discuss considerations of confidentiality.) If this is a commercially sponsored project, it will generally be provided by the sponsors. If you need assistance in any aspect of study design contact the Clinical Research Department at 208-2238 for a research consultation.
- Investigator's Brochure if Drug Study
- Patient Recruitment Advertisements if any
- Informed consent document
- Questionnaire or Data Collection Form
- Curriculum Vitae (CV) of the investigators (unless previously submitted)
- Completion of CITI training
- If any investigators are active duty military, please contact the CRC to determine additional requirements.

Research cannot begin until approval by the MVH Human Investigation and Research Committee (HIRC) has been received.