IRB Content Webpage

# Introduction page

# Policy and procedures

## Policy # 1& 2

## SOPs

# IRB Meeting Schedule and dateline

# Regulations & resources

## Helsinki

## Belmont

## Health & Human Services (HHS)

## Common Rule

## FDA

## NIH Decisional Charts

# Forms & templates

## IRB Application

## Supplements

### Supplement A- Waiver Application

### Supplement B- International Research

### Supplement C- Research Involving Children

### Supplement D- Cognitive Impaired

### Supplement E- Stored data

### Supplement F- Research involving drugs or biologicals

## IRB Checklist

## Informed Consent Template

### Instructions and template of the Informed Consent- English version

### Template Informed Consent- Spanish version

### Template of the Broad Consent- English

## Financial conflict of interesting form- FCOI

## Translation Attestation Form

# IRB Committee

## Members of the IRB

## IRB Member’s responsibility

## Access to members only- to post the research studies and all documents to be evaluated and merge the members’ evaluation in one document.

### Research Evaluation form

# Education and trainings

## CITI electronic address

# Glossary