Forms & Consent Templates

Consent Templates

The IRB recommends the use of the consent templates to help researchers meet the legal requirements for consent.

See the Informed Consent Process page [1] for more information about the consent process.

Medical(SoM) [2]

- School of Medicine (SoM)
- Lucile Packard Children's Hospital (LPCH)
- Stanford Hospital and Clinics (SHC)
- Veteran's Affairs (VA) Hospital
- Psychology fMRI studies

Social & Behavioral Research (Non-Medical) [3]

- Business
- Education
- Engineering
- Humanities & Sciences
- Law

Checklists

The IRB uses these checklists and forms to review protocols for compliance with regulations, policies and guidance:

Staff Checklists

Exemption Eligibility [4]

Protocol Checklists:

- Protocol - Medical [5]
- Protocol - Expedited (initial review) [6]
- Protocol - Chart Review (initial review) [7]
- Protocol - Nonmedical [8]
- Protocol - sIRB Checklist [9]
  Research Involving VA Studies [10]; see Reviewing Veterans Affairs (VA) Research [11] for additional requirements
- Exemption from IRB Review: Emergency Use of a Test Article [12]
• Single Patient IND/IDE [13]

Other Federal Agency Requirements:
• Dept. of Defense (DoD) [14]
• Dept. of Education (ED) [15]
• Dept. of Energy (DOE) [16]
• Dept. of Justice (DOJ) [17]
• Environmental Protection Agency (EPA) [18]

Informed Consent:
• Informed Consent [19](medical: clinical studies)
• Informed Consent [20](medical: expedited/minimal risk)
• Informed Consent [21](nonmedical: surveys, social, behavioral, education research)
  Neonates [22]

Continuing Review:
Continuing Review - FULL/EXPEDITED [23]

Reviewer Checklists:
• Medical [24]
• Nonmedical [25]

Source URL: https://researchcompliance.stanford.edu/panels/hs/forms/forms-templates

Links
[4] https://stanfordmedicine.box.com/shared/static/z5hek9y2jsf91oa5awurgxqb9fyv44qo.docx
[5] https://stanfordmedicine.box.com/shared/static/e50dlny0g5rzlmskm2dbvp17swngi0t.docx
[8] https://stanfordmedicine.box.com/shared/static/tt7fwv0uxws4uwc0k52f47k9z5buq0u.docx
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