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INSTITUTIONAL REVIEW BOARD

IRB CHECKLIST FOR INITIAL SUBMISSION (FORM 2.2)

Protocol package for clinical trial and/or sponsor-initiated studies:

- Letter of Application & Full protocol
- Executive summary that follows research project proposal format
- Investigator's Brochure
- Data collection form/s
- Informed Consent form (English, Tagalog, and local dialect (Hiligaynon))
- Budget
- Curriculum Vitae of the Principal Investigator and his/her co-investigators
- GCP Certificate of the Principal Investigator and his/her co-investigators
- Declaration of No Conflict of Interest of Investigators/Researchers
- Photocopy of valid PRC Licence
- GANTT Chart (as necessary)
- Advertisement

Protocol package for researcher-initiated studies (residents, staff, students, government sector and private sector, etc.):

- Letter of Application & Full protocol (Chapter 1-3)
- Executive summary that follows research project proposal format
- Data collection form/s
- Informed Consent form (English/Tagalog, and local dialect (Hiligaynon))
- Budget (as necessary)
- Curriculum Vitae of the PI and co-investigators
- Declaration of No Conflict of Interest of Investigators/Researchers
- GCP Certificate (for intervention studies only)
- Photocopy of valid PRC Licence
- GANTT Chart (as necessary)
- Certificate of Technical Review