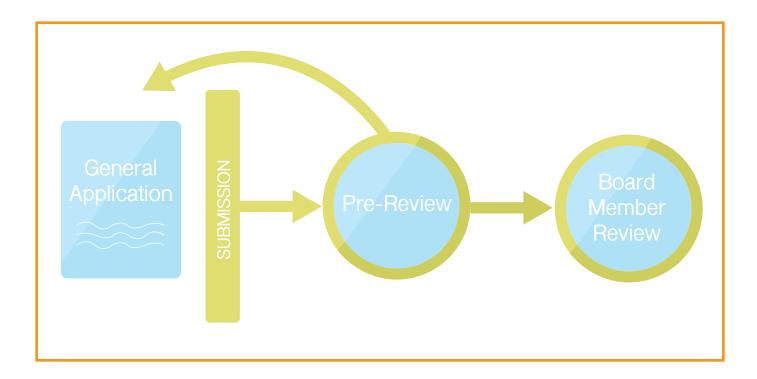
Pre-Review Rubric

When your General application is first submitted through Protis, the IRB Staff Assistant conducts a pre-review to check for completeness. If the application is incomplete, it will be returned to you for revision prior to review. On the next page of this document, you will find the rubric used during this stage of the process.

Please refer to the rubric as you fill out your application to secure a faster review time.



Return to PI if 1 or more are checked.

Return to PI if 3 or more are checked.

•	Basic Project Information		Members of the research staff don't have current CITI certification.
	The PI does not have current CITI training A student researcher is listed as the PI.		(For student projects) The signed committee cover page hasn't been
•	Students, Dates & Funding	Н	uploaded (For student projects) The faculty
•	Scientific Validity		advisor hasn't completed and uploaded the SVRC.
•	Project Details Participant Population		There are inconsistencies between the research proposal and the information in the protocol.
	A participant population is missing. Be sure to identify all the groups participating and to respond to protocol prompts for each group. For example, if you are collecting data from children and teachers, you should address privacy for both.	Ē	How researchers will invite individuals to participate is unclear. Recruitment materials (verbal scripts,
•	Vulnerability of Participants		email invitations, flyers, SONA post language, etc.) are missing.
	Participants include vulnerable populations, but none are marked in the vulnerability tab.		Applicable vulnerabilities haven't been checked.
•	Informed Consent		
	The PI used an outdated template to develop the informed consent documents. The newest version of the template is on the website and is linked within the protocol.		
	A consent form is missing for a population involved in the research.	П	A step-by-step description of study
	A HIPAA Authorization Form is missing. (For research involving the collection of private health information protected by HIPAA)	Н	procedures isn't included.
	Procedures	ы	Some measures/data collection sheets are missing.
	A majority of the data collection instruments are missing.		Risks and benefits are inconsistent with the information in the Informed Consent document(s).
•	Risks & Benefits		
•	Privacy and Confidentiality		Researchers have incorrectly distinguished between confidentiality and privacy.
	Participants will be identifiable to the researchers (whether by in-person interaction or through collected personal information), but the researcher selected that participants are not identifiable.		The information and time lines in the Privacy and Confidentiality tab are inconsistent with information in the Informed Consent document(s).
	Reporting, Health, COI, Assurance, & Submit tabs		