

INSTITUTIONAL REVIEW BOARD CHECKLIST

Submitted By: _____

Date: _____

As part of the IRB process, the Principal Investigator must supply the following required information and documentation listed below. **Include completed checklist with your submission.**

| | Brief description of the background and purpose of the research. Spell out | |
|--|---|---|
| _ | | rms the first time they are used; and use consistent terminology. |
| | Data C | collection instruments (surveys, interview questions, etc.), or |
| | | Not applicable as research involves no test items. |
| Description of the procedure for ensuring the following for research participants: | | |
| | | Anonymity (the researcher will not ever be able to link names or personal identifiable information), <u>or</u> |
| | | Confidentiality (the researcher will be able to link names to the data but will keep that information private), <u>or</u> |
| | | Neither anonymity nor confidentiality |
| For online surveys only , indicate how your data are stored: | | |
| | | Online survey data are emailed to the researcher |
| | | Data are stored online |
| | | A combination of the above. (Please specify in your procedural section) |
| | | Physical location (e.g., office) |
| | | Other (please specify) |
| | Consent document(s) and script for obtaining informed consent. | |
| | Copy of recruitment materials, if applicable (e.g., emails recruiting participants and/or the description of the study in online participant pool sign-up systems). | |
| | | How and when participant eligibility criteria are communicated (if |

- How and when participant eligibility criteria are communicated (if necessary).
- Letters of IRB approval from cooperating institutions (if applicable), or
 - \Box Not applicable as no other institutions are involved.
- Evidence of human subjects' protection training completed within two years

Principal Investigator's Signature