

## **Exemption Category 3**

Research involving benign behavioral interventions in conjunction with the collection of information from an adult subject through verbal or written responses (including data entry or audiovisual recoding if the subject prospectively agrees to the intervention and information collection and at least one of the following criteria is met:

Data collected anonymously	No Risk if inadvertently disclosed	Sensitive identifiable data*
(A) The information obtained is rec- orded by the investigator in such a manner that the identity of the hu- man subjects cannot readily be ascer- tained, directly or through identifiers linked to the subjects.	(B) Any disclosure of the human sub- jects' responses outside the research would not reasonably place the sub- jects at risk of criminal or civil liability or be damaging to the subjects' finan- cial standing, employability, educa- tional advancement, or reputation.	(C) The information obtained is rec- orded by the investigator in such a manner that the identity of the hu- man subjects can readily be ascer- tained, directly or through identifiers linked to the subjects. *Limited IRB Review Required

**Benign Behavioral Intervention:** Brief in duration, harmless, painless, not physically invasive, not likely to have a significant adverse lasting impact on the subjects, and the investigator has no reason to think the subjects will find the interventions offensive or embarrassing.

	Studies not eligible for Exempt #3 include:	Deception or Incomplete Disclosure	
*	Children and/or vulnerable populations (e.g. decision- ally-impaired).	Exemption only if all of the following are met: Deception/incomplete disclosure are authorized by the subject—prospectively informed that they may be una-	
•	Collecting inherently high risk data, such as HIV status, criminal behaviors, suicide, medical information, etc.	ware of or misled regarding the nature/purpose of the study.	
•	Physical procedures: Blood Pressure, EEG, activity trackers, eye trackers, blood draws, ultrasound, etc.	The information withheld is unlikely to meaningfully effect willingness to participate.	
* *	esting medical devices. inking study data to other data such as medical rec- rds, student records, administrative data.	The deception/incomplete disclosure does not preclude disclosure of risks or discomfort, or voluntary participa- tion.	
•	Incomplete disclosure or deception that is not pro- spectively disclosed.	The deception/incomplete disclosure does not involve misleading subjects about anything that may be substan- tively upsetting or cause distress.	
	Examples: May be eligible for Exempt #3:	Examples: May <i>not</i> be eligible for Exempt #3:	
•	Health eating intervention with a pre/post question- naires. Study among young adults evaluating preferred snack foods following viewing a movie. Participants may watch a happy movie and others may watch a sad movie. Movie selection is manipulated based on pre- survey data. Participants are told in the consent form that they will be misled or not fully informed about the nature/purpose of the study.	<ul> <li>Healthy eating intervention with pre/post question- naires and use of student academic record and food buying habits collected from student ID purchases.</li> <li>Study among young adults evaluating preferred snack foods following viewing a movie. Participants may watch a happy movie and others may watch a sad movie. Movie selection is manipulated based on pre- survey data. Participants are unaware that the move selection is manipulated or the full nature/purpose of the study.</li> </ul>	