Exemption Category	Exemption Description	Conditions/Allowances/Limitations
1	Research in established or commonly accepted education settings that involves normal educational practices	Not likely to adversely impact students' opportunity to learn or assessment of educators
2	Research only includes educational tests, surveys, interview, public observations if at least one of the following criteria is met:	Data collection only; May include visual or auditory recording; May **NOT** include Intervention
	(i) Recorded information cannot readily identify the subject (directly or indirectly linked)	Surveys & interviews: no children Educational tests or observation of public behavior can only include children when investigators do not participate in activities being observed
	(ii) Any disclosure of responses outside of the research would NOT reasonably place subject at risk (criminal, civil liability, financial, employability, educational advancement, reputation).	Surveys & interviews: no children Educational tests or observation of public behavior can only include children when investigators do not participate in activities being observed
	(iii) Information is recorded with identifiers & IRB conduct Limited Review	No children
3	Research involving Benign Behavioral Interventions through verbal, written responses, (including data entry or audiovisual recording**) from adult subject who prospectively agree and ONE of following met:** A. Recorded information cannot readily identify the subject (directly or indirectly/linked) B. Any disclosure of responses outside of the research would NOT reasonably place subject at risk (criminal, civil liability financial, employability, education advancement, reputation) C. Information is recorded with identifiers & IRB conducts Limited Review	No children; Many not include medical interventions; Subject prospectively agrees; Benign Behavioral Intervention must be: • Brief in duration • Painless/harmless • Not physical invasive • Not likely to have a significant adverse lasting impact • Unlikely that subjects will find interventions offensive or embarrassing No deception unless participant prospectively agrees

4	Secondary research for which consent is not required: use of identifiable information or identifiable biospecimen that have been or will be collected for some other 'primary' or 'initial' activity, if ONE of following criteria met:	No primary collection from subjects for the research; Allows both retrospective and prospective secondary use.
	(i) Biospecimens or information is publically available	Must be publically available
	(ii) Information recorded so subject cannot readily be identified (directly or indirectly/linked); Investigator does not contact subjects and will not re-identify the subjects	
	(iii) Collection and analysis involving investigators use of Identifiable Health Information when use is regulated by HIPAA 'health care operations" or "research" or "public health activities and purposes"	HIPAA still applies HIPAA protections include authorization or waiver of authorization;
	(iv) Research information collected by or on behalf of federal government using government generated or collected information obtained for non-research activities	If research generates identifiable private information it is subject to specified federal privacy laws
5	Research and demonstration projects supported by aa Federal Agency/Dept. AND designed to study public benefit or service programs	Must be posted on a federal web site
6	Taste and Food Qualty and Consumer Acceptance Studies	

EXEMPTION 7 AND 8 WILL NOT BE IMPLEMENTED AT WHITWORTH

7	Storage or maintenance of identifiable private information or identifiable biospecimens for secondary research for which broad consent if required	IRB may waive consent requirement; Of; All requirement for Broad Consent must be met and refusals to consent must be tracked; the IRB may not waive consent for use of identifiable material for any individual who refuses
8	Secondary research involving use of identifiable private information or identifiable biospecimens for which Broad Consent was required	Privacy and confidentiality protections adequate; Broad Consent was obtained; Documented or documentation waived Return of research results not allowed; Refusal s to consent must be tracked; The IRB may not waive consent for use of identifiable material for any individual who refuses