

IRB – Child/Teen Assent Form (Ages 7-18)		
For all studies		
Title of Research Project:		
Principle Investigator(s):		
Purpose of Research: In age-appropriate language, explain why the study is being done.		
Procedures and Location: Describe what the participant will do, how often, and where, using plain, easy to understand language.		
Duration of Participation: Explain how long participation will last.		
Risks or Discomforts: Easily explain any risks, e.g., "You might feel shy answering questions.".		



Benefits: Describe any direct or indirect benefits in clear, understandable terms.		
Alternatives (if applicable): Explain what the participant can do instead of participating.		
Compensation or Incentives (if any): State what the participant will receive.		
Confidentiality: Explain how information will be kept private and who can access it, in terms that will make sense to the child or adolescent.		
Voluntary Participation and Right to Withdraw: Explain that participating is voluntary, and the he/she can stop at any time, using age-appropriate language.	at	



Pl's Contact Information Regarding Research Questions:	
Contact for the Office of Sponsored Programs and Researc	h:
Operation Observan	
Caroline Stanton	
Compliance and Research Officer	
Office of Sponsored Programs and Research	
205-652-3403, cstanton@uwa.edu	
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Notices (if applicable): Easily describe any special informati	•
as whether audio or video recordings will be made, data wi	
to participants, or if biospecimens could result in commerci	ai pront.
Broad Consent for Future Use (if applicable): Are you willing	ng to give consent for future
research use of de-identifiable information or biospecimen	
☐ Yes ☐ No	
Child's/Teen's Assent & Signature:	
☐ Yes, I want to be in the study.	
☐ No, I do not want to be in the study.	
Printed Name of Child/Teen:	
	
Signature of Child/Teen (if appropriate):	Date:
Printed Name of Researcher Obtaining Consent:	
	
Signature of Researcher Obtaining Consent:	Date: