

## Checklist for the Informed Consent (or Assent) Process

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*This checklist helps to assure that all components of the consent process are addressed, and in the proper order. Adapt to your own study as needed*

Study Name: \_\_\_\_\_ Participant ID: \_\_\_\_\_

Items Addressed	Comments
<input type="checkbox"/> List persons present during the informed consent process and the consent signee.	Persons Present:
<input type="checkbox"/> List the person who explained the details of the study participation.	Name:
<input type="checkbox"/> Discussed purpose of research and procedures.	
<input type="checkbox"/> Discussed risks and benefits of study participation.	Questions asked and answered:
<input type="checkbox"/> Discussed alternatives to research.	
<input type="checkbox"/> Discussed that participation is voluntary and participants may withdraw at any time.	
<input type="checkbox"/> Discussed issues of confidentiality.	
<input type="checkbox"/> Discussed potential study-associated costs.	
<input type="checkbox"/> Assessed if family appears to understand all terms of participation and agree to enrollment. (parent <u>and</u> child if assent is needed)	Demonstrated through: <input type="checkbox"/> Talk back method <input type="checkbox"/> Q&A <input type="checkbox"/> Other <hr style="width: 20%; margin-left: 0;"/>
<input type="checkbox"/> The consent document was signed prior to the performance of any study-related procedures.	
<input type="checkbox"/> A signed copy of the consent was provided to the participant, including investigator/research team contact information.	

Investigator or IRB-approved delegate who conducted the consent process and completed this form.

Signature:	Date:	
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