Informed Consent Form

[Title of your study]

Principal investigator: [Name]
RWTH Aachen University
Email: [Your RWTH email address]

Purpose: [Explain the goal of your study. This should communicate what your intentions are and justify collecting data from participant. Be careful not to prime the participant, e.g., by letting her know you are looking for a particular result.]

Procedure: [Explain the various steps in your study, what tasks you expect the participant to do (Should she think aloud? Can she ask you for help?), and any other considerations the participant should be aware of before the study begins. Also indicate what information you will capture in the study (video, audio, demographic information, screen capture, etc.).]

Risks: [Describe possible risks in partaking the study, e.g., mental or physical fatigue, duration of the study, and the procedure for possible breaks or an early termination due to distress during the study.]

Confidentiality: [Explain if and how the participant' information will be kept confidential. If you need to divulge private information to publish your study findings, you need to get the participant's consent.]

Costs and Compensations: [Does participation in your study involve a monetary cost to the participant? Is the participant compensated in some way for her time, e.g., drinks, snacks, or monetary rewards.]

☐ I have read and understood the information on this form.

☐ I have had the information on this form explained to me.

Participant's Name	Participant's Signature	Date
	Principal Investigator	Date