UNIVERSITY OF CALIFORNIA, SAN DIEGO
CONSENT TO PARTICIPATE IN RESEARCH

1. Study Title and Number

Title: Naturalistic Stimuli of Person Perception
Study: #810260

2. Principal Investigator

Chujun Lin, Assistant Professor, Department of Psychology

3. Principal Investigator Phone Number, Research Team Number, and Emergency Contact Number

(626)-689-2886

4. Study Overview

This research study is being conducted to generate a new database of images, audios, and videos of people and their interactions with others in daily life to facilitate research on understanding how people perceive each other.

We are inviting you to participate in a research study because you are aged 18 and older, with normal or corrected-to-normal vision and hearing, fluent in English, and with at least high school education.

This form explains the research so that you may make an informed decision about participating.

- Research is voluntary - whether or not you participate is your decision. You can discuss your decision with others (such as family, friends or another physician).
- You can say yes, but change your mind later.
- If you say no, we will not hold your decision against you.
- You can say no even if the person inviting you is part of your healthcare team.
- Your decision will not affect your health care or other benefits you may be entitled to.
- Please ask the study doctor or study team questions about anything that is not clear, and feel free to ask questions and mention concerns before, during, and after the research.
- You may consult with friends, family, a personal doctor, or anyone else before deciding whether or not to be in the study.
- You will be given a copy of this consent form and the Participant's Bill of Rights.

The purpose of this research study is to collect images, audio recordings, or video recordings about you in daily life to use as stimuli to show other participants for scientific research purposes. You have complete control over the content of the stimuli (which should be appropriate) and whether these stimuli can be stored and shared (you will be given a data release survey at the end of the study).

You will first read the informed consent and instructions of the study. The instructions will let you know which type of stimuli are being collected and the requirements of these stimuli. If you decide to proceed, you will either participate via our online studies where you will upload the stimuli of your choice by yourself or via our in-person studies where our researchers will record the stimuli of you. After providing stimuli, you may also be asked to fill out questionnaires about yourselves such as your demographics, moods, personalities, and attitudes, and questionnaires about other participants based on their stimuli such as inferring their demographics, moods, personalities, and attitudes. At the end of the study, you will be asked to fill out a data release survey, during which you can decide whether to completely erase your data or allow the storage and/or share of your identifiable stimuli or their de-identified derivatives. The study will last up to 1 hour.

The most common risks or discomforts of this study include psychological, social, and reputational risks. However, these risks are no more than those that would occur in any everyday internet use or those that would occur in any everyday conversation with another person. The multi-tiered data release survey mentioned above will give you complete control over what data will be retained, and what - if any - will be shared. If you are participating in our in-person studies, our research assistants will also supervise the interactions in real time to ensure your safety and well-being.

We cannot promise any benefit to you or to others from you participating in this research. However, possible benefits include gaining knowledge of scientific research processes and having the opportunity to engage with others in the local community if you participate in our in-person interaction studies.

The alternative to being in this study is not to participate.

More detailed information about this research study is provided below.

5. Whom can I talk to if I have questions?

If during your participation in the study you have questions or concerns, or if you think the research has hurt you, contact the research team at the numbers listed in Section 3 on the first page of this form. You should not agree to participate in this study until the research team has answered any questions you have about the study, including information contained in this form.

If before or during your participation in the study you have questions about your rights as a research participant, or you want to talk to someone outside the research team, please contact:

- UC San Diego Office of IRB Administration at 858-246-4777 or irb@ucsd.edu

6. How many people will take part?

We plan to study 500 people here. The research will include online participants recruited around the country and beyond, as well as in-person participants recruited in the California regions.

7. What happens if I take part in the research?

Here is what will happen to you if you agree to be in this study:

- Images, audios, or videos of you will be collected;
- Survey responses about your demographics, moods, personalities, and attitudes will be collected;
- Survey responses about your inferences of others' demographics, moods, personalities, and attitudes will be collected based on others' stimuli - there are no right or wrong answer so please provide your most honest responses;
- Survey responses about your preferences for the storage and share of your stimuli will be collected.

As you read this form, ask questions if something is not clear.

8. What are the risks and possible discomforts?

Participation in this study may involve risks or discomforts. We anticipate minimal psychological, social, and reputational risks that are no more than those that would occur in any everyday internet use or those that would occur in any everyday conversation with another person. The multi-tiered data release survey will give you complete control over what data will be retained, and what - if any - will be shared. Research assistants

will also supervise in-person interactions in real time to ensure the safety and well-being of participants as they interact.

9. How will information about me be protected?

While we cannot guarantee complete confidentiality, we will limit access to information about you. Only people who have a need to review your information, documents, or specimens will have access. These people might include:

- Members of the research team and other staff or representatives of UCSD whose work is related to the research or to protecting your rights and safety.
- Representatives of the study sponsor or product manufacturer
- Representatives of Federal and other regulatory agencies who make sure the study is done properly and that your rights and safety are protected.

As a part of this study, photographs and/or videos will be taken of your face and/or parts of your body, and/or audios will be taken of your voice. These stimuli will be subject to the same confidentiality conditions described above. Even so, someone who knows you well, may be able to identify you from these stimuli and know you are participating in this study.

To minimize this risk, we will take the following precautions: We will provide you with a data release survey at the end of the study, which will allow you to decide what stimuli of you to be erased, stored, and shared. The stimuli that you agree to be stored will be securely stored with password protection and regular backups. The stimuli that you agree to be shared will be shared only for scientific research purposes and only if a data request form is filled out by the party wishing to use the stimuli.

The results of this study may be published once the study is completed. However, we will keep your name and other identifying information confidential. We expect this study will be completed in 3 years. This is only an estimate and the actual time to complete the study may be longer or shorter depending on a number of factors.

Under California law, we must report information about known or reasonably suspected incidents of abuse or neglect of a child, dependent adult or elder including physical, sexual, emotional, and financial abuse or neglect.

10. Will I need to pay to participate in the research?

There will be no cost to you for participating in this study.

11. What if I agree to participate, but change my mind later?

You can stop participating at any time for any reason, and it will not be held against you. Your choice will not affect any treatment relationship you have with healthcare providers at UC San Diego Health or any services you receive from them. No matter what you decide, there will be no penalty to you. You will not lose medical care or any legal rights.

12. What will happen to information collected from me?

The data we collect with your identifiable information (your images, audios, and videos) as a part of this study may be used to answer other research questions or may be shared with other investigators for other research, depending on your preferences indicated in the data release survey at the end of the study. Your name and

contact information provided in this informed consent will never be shared and will be stored separated from your stimuli.

13. Will I be compensated for participating in the research?

If you agree to take part in this research, we will provide you $10 for your time and effort. This compensation is based on a participation time of 20 minutes.

14. What else is important for me to know?

You will not be provided any clinically relevant information that may pertain to your health. You will not be provided a summary of the research findings.

15. What are my rights when providing electronic consent?

California law provides specific rights when you are asked to provide electronic consent:

- You have the right to obtain a copy of the consent document in a non-electronic format.
- You have the right to provide consent in a non-electronic format.
- If you change your mind about electronic consent, you have the right to request your electronic consent to be withdrawn and you can then provide consent in a non-electronic format; however, a copy of your electronic consent will be maintained for regulatory purposes. If you wish to withdraw your electronic consent please tell the study team.

This agreement for electronic consent applies only to your consent to participate in this research study.

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CONSENT TO PARTICIPATE IN RESEARCH

Participant

I have received a copy of this consent document and a copy of the "Experimental Participant's Bill of Rights" to keep. I agree to participate in the research described in this form.

Printed Name of Participant

Signature of Participant Date

Person Obtaining Consent

I document that:

- *I (or another member of the research team) have fully explained this research to the participant.*
- *I have personally evaluated the participant's understanding of the research and obtained their voluntary agreement.*

Printed Name of Person Obtaining Consent

Signature of Person Obtaining Consent Date

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Experimental Participant's Bill of Rights

Every individual asked to participate in a research study has the right to be:

1. Informed about the nature and purpose of the study.

2. Provided an explanation of the procedures to be followed in the research study, and whether any of the drugs, devices, or procedures is different from what would be used in standard practice.

3. Given a description of any side effects, discomforts, or risks that you can reasonably expect to occur during the study.

4. Informed about any benefits that would reasonably be expected from the participation in the study, if applicable.

5. Informed about of any alternative procedures, drugs, or devices that might be helpful, and their risks and benefits compared to the proposed procedures, drugs or devices.

6. Told of the types of medical treatment, if any, available if complications should arise.

7. Provided an opportunity to ask any questions concerning the research study both before agreeing to participate and at any time during the course of the study.

8. Informed that individuals can refuse to participate in the research study. Participation is voluntary. Research participants may refuse to answer any question or discontinue their involvement at any time without penalty or loss of benefits to which they might otherwise be entitled. Their decision will not affect their right to receive the care they would receive if they were not in the experiment.

9. Provided a copy of the signed and dated written consent form and a copy of this form.

10. Given the opportunity to freely decide whether or not to consent to the research study without any force, coercion, or undue influence.

If you have any concerns or questions regarding the research study contact the researchers listed at the top of the consent form.

If you are unable to reach a member of the research team and have general questions, or you have concerns or complaints about the research study, research team, or questions about your rights as a research participant, please contact:

- UC San Diego Office of IRB Administration at irb@ucsd.edu or 858-246-4777