

**STANFORD UNIVERSITY Research Information Sheet**

Protocol Director: Tracey McLaughlin, MD, and  
Christopher Gardner, PhD

*IRB Use Only*

Approval Date: March 30, 2018  
Expiration Date: February 28, 2019

Protocol Title: **Common consent for Stanford diabetes clinical research registry**

**ONLINE SCREENING QUESTIONNAIRE  
WAIVER OF DOCUMENTATION**

**Please acknowledge one of the following:**

**- You are an adult participant in this study.**

**OR**

**-You are the parent or guardian granting permission for a child in this study.**

**The following information applies to the adult participant or to the child or ward. If the participant is a child or ward, the use of "you" refers to "your child" or "your ward."**

**DESCRIPTION:** Thank you for your interest in participating in a diabetes-related research study. The purpose of this screening questionnaire is to create a clinical research registry from which investigators within the Stanford Diabetes Research Center can recruit participants. In order to find out whether you are eligible for particular studies, you will be asked to complete a pre-screening questionnaire. There are two steps involved in this pre-screening process: 1) consent to be screened via the questionnaire, and 2) if you agree to the consent, complete the online screening questionnaire to determine whether you are eligible to attend an in-person screening visit with one or more groups part of the Stanford Diabetes Research Center in the near or distant future. Your information will be kept as confidential as possible as required by law. This survey is encrypted for additional security.

**TIME INVOLVEMENT:** Your participation will take approximately 10 minutes to complete the questionnaire.

**RISKS AND BENEFITS:** There are no foreseeable risks associated with this pre-screening survey. You may benefit by becoming eligible for one or more diabetes-related studies conducted at Stanford University. **We cannot and do not guarantee or promise that you will receive any benefits from this study.**

**PAYMENTS:** You will not receive payment for your participation.

**PARTICIPANT'S RIGHTS:** If you have read this form and have decided to participate in this project, please understand your participation is voluntary and

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you have the right to withdraw your consent or discontinue participation at any time without penalty or loss of benefits to which you are otherwise entitled. The alternative is not to participate. You have the right to refuse to answer particular questions. With your permission, your identity will be made known to research groups within the Stanford Diabetes Research Center for recruitment purposes only.

### **CONTACT INFORMATION:**

*Questions:* If you have any questions, concerns or complaints about this research, its procedures, risks and benefits, contact the Protocol Director, Tracey McLaughlin, MD at 650-721-1300.

*Independent Contact:* If you are not satisfied with how this study is being conducted, or if you have any concerns, complaints, or general questions about the research or your rights as a participant, please contact the Stanford Institutional Review Board (IRB) to speak to someone independent of the research team at (650)-723-5244 or toll free at 1-866-680-2906. You can also write to the Stanford IRB, Stanford University, 3000 El Camino Real, Five Palo Alto Square, 4th Floor, Palo Alto, CA 94306.

### **Please print a copy of this page for your records**

After reading the above information, do you consent to continuing with the screening questions? **If you agree to participate in this research, please** click "yes" below and continue to the next page and start Step 2, the online screening questionnaire.