Content for Research Information Sheet
Overdose Prevention in Primary Care

Description of purpose of the study

- This is a University of Washington research study funded by the Centers for Disease Control and Prevention as one of four projects supported by the Harborview Injury Prevention Research Center.
- Identify barriers and facilitators of implementing harm reduction practices to prevent overdose among primary care patients.

For Interviews:

- Description of procedures
  - Brief 5-minute survey to provide demographic and practice information via REDCap
  - Interviews with clinicians (n = 20) will take ~45-60 minutes.
  - Interviews will be audio and video recorded. We will ask permission prior to starting the recording.
  - Interview questions will explore barriers and facilitators to integrating harm reduction practices, existing strategies to prevent overdose, implementation strategies used, and potential new solutions.
  - Participation in co-design activities is optional.

- Description of risks and benefits
  - Risks: Potential discomfort discussing harm reduction, discussing clinic barriers (e.g., disclosure of colleagues' stigma), and addressing challenges faced by patients who use drugs
  - Benefits: Contributing to improved overdose prevention practices primary care clinics

- Description of confidentiality protection (e.g., data privacy and security)
  - The information and/or specimens that we obtain from you for this study might be used for
  - future studies. We may remove anything that might identify you from the information and specimens. If we do so, that information and specimens may then be used for future research studies or given to another investigator without getting additional permission from you. It is also possible that in the future we may want to use or share study information that might identify you. If we do, the Harborview Injury Prevention Research Center ethics consultant will decide whether or not we need to get additional permission from you. If you are a resident, we will not provide any information about your participation to supervisors. Your participation is voluntary and will not be used in any evaluative capacity.

- Description of compensation for participation
  - $100 Tango card for interviews and brief survey

- Study withdrawal information
  - Participation is voluntary, and you can withdraw at any time or refuse any question/activity.

We have a Certificate of Confidentiality from the United States National Institutes of Health. These protections only apply to data held in the United States.

This helps up protect your privacy. The certificate means that we do not have to give out information, documents, or samples that could identify you even if we are asked to by a court of law in the United States. We will use the Certificate to resist any demands for identifying information.

We can't use the Certificate to withhold your research information if you give your written consent to give it to an insurer, employer, or other person. Also, you or a member of your family can share information about yourself or your part in this research if you wish.

There are some limits to this protection. We will voluntarily provide the information to:

- a member of the United States government who needs it in order to audit or evaluate the research;
- individuals at the institution(s) conducting the research, the funding agency, and other groups involved in the research, if they need the information to make sure the research is being done correctly;
- individuals who want to conduct secondary research if allowed by federal regulations and according to your consent for future research use as described in this form;
- to relevant authorities as required by other Federal, State, or local laws.

The Certificate expires when the CDC funding for this study ends. Currently this is July 30, 2030. Any data collected after expiration is not protected as described above. Data collected prior to expiration will continue to be protected.

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