IRB #: <u>xxxx</u>

IRB Approval Date: xxxx

| Study title | The test of the kiosk prototype |
|-------------|---|
| Researcher | Yuzhou Wu, Linyi Gao, Yongliang Tan, Sicheng Wang |

We're inviting you to participate in a test. Participation is completely voluntary. If you agree to participate now, you can always change your mind later. There are no negative consequences, whatever you decide.

What is the purpose of this study?

We want to test if the functionality of our kiosk can work properly.

What will I do?

In our test, you'll do the implementation guided by our moderator.

Risks:

The personal information may be disclosed. To reduce the risk, we will use de-identified name during the test.

Other Study Information

| Estimated number of participants | 100 non-disabled people and 10 disabled people | |
|----------------------------------|--|--|
| How long will it take? | About 10 minutes | |
| Costs | None | |
| Future research | Your result won't be used or shared for any future research studies. | |

What if I am harmed because I was in this study?

If you're harmed from being in this study, let us know. If it's an emergency, get help from 911 or your doctor right away and tell us afterward. We can help you find resources if you need psychological help. You or your insurance will have to pay for all costs of any treatment you need.

Confidentiality and Data Security

We'll collect the following identifying information for the research: your name, gender and age. This information is necessary because we wish each kind of people can use our Kiosk smoothly.

| Where will data be stored? | On our computers |
|----------------------------|------------------|
| How long will it be kept? | Three months |

| Who can see my data? | Why? | Type of data |
|--|---------------------------------|---------------------|
| The researchers | To conduct the test and analyze | The records of user |
| | the data | implementations |
| The IRB (Institutional Review Board) | To ensure we're following laws | The records of user |
| The Office for Human Research Protections (OHRP) or other federal agencies | and ethical guidelines | implementations |

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Contact information:

| For questions about the research | Yuzhou Wu | XXX-XXX-XXXX |
|---|---|--------------|
| | Linyi Gao | |
| | Sicheng Wang | |
| | YongLiang Tan | |
| For questions about your rights as a research participant | IRB (Institutional Review Board; provides ethics oversight) | XXX-XXX-XXXX |
| For complaints or problems | Yuzhou Wu Linyi Gao Sicheng Wang YongLiang Tan | XXX-XXX-XXXX |
| | IRB | XXX-XXX-XXXX |

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| • | | in this study, sign on the lines below. withdraw from the study at any time. |
| Name of Participant (print) | | menanan mem ene seday de dify clinici |
| . " | | |
| Signature of Participant | | Date |