

## On-line Video Informed Consent Research Project Survey

Dear volunteers

Thank you for your time to participate this research project.

This survey is being conducted in support of the on-line video informed consent study. The purpose of this survey is to evaluate the understanding of the whole one-on-one on-line video informed consent procedure. Data collected from this survey will make a better understanding of this project, and provide an evidence of on-line video informed consent in future as reference..

Your participation in this research project is completely voluntary. Your participant privacy and confidentiality will be protected throughout this study. All documentation will be kept strictly confidential. Information from the survey will be coded to preserve participant anonymity and confidentiality. Data collected in support of this research project will be summarized, in anonymous format, in the body of the final report for publication.

Thank you again for your support and cooperation in this survey.

Department of Pharmacology,

Aerospace center hospital

26<sup>th</sup>, March, 2020

### Section 1. General Information

1. Gender    male         female
2. Age        18-25         26-30         31-45          $\geq$ 46

### Section 2. Evaluation of video informed consent process

1. Have you participated in any phase I clinical trials before ?    Yes     No
2. Were you satisfied with doctors' explanation in this video informed process?  
Yes     No
3. Did you fully understand what the doctor explained?  
Yes, fully         Yes         Yes, basically         No
4. Do you think the effectiveness of video-based informed process is consistent with the onsite process?  
Yes         Yes, basically         No
5. Is the doctor's explanation of the informed consent form easy to understand? Yes     No

6. What points of information did you focus on through this video informed consent process?  
 Risk  Drug information  Blood volume  Period  Adverse reaction   
 Privacy  Compensation
7. Have the doctor fully explained the information what you are concerned? Yes  No
8. What is your overall evaluation of the video-based informed process?  
 Very good  Good  OK  Bad
9. Are you willing to participate the video-based informed process again? Yes  No
10. What are the advantages of video-based informed method?  
 Save time  Save transportation cost  Avoid gathering  Graphical form   
 None  Others \_\_\_\_\_
11. What are the drawbacks of video-based informed method?  
 Unstable network  Noisy environment  Fast speed  Network flow limit   
 None  Others \_\_\_\_\_

### **Section 3. Trial related information of informed consent form**

**Please truthfully answer the questions about this trial, if you can't remember them clearly, the doctor would explain again in the face-to-face in person discussion.**

1. Are you able to answer the time, schedule and duration of this trial?  
 Yes , please go on filling: the admission date \_\_\_\_\_ and the duration days \_\_\_\_\_  
 No
2. Are you able to answer the indication of IMP?  
 Epilepsy  Hypotension  Parkinson's disease  Cerebral infarction
3. Are you able to answer the times of oral drug administration during the trial?  
 Once  Twice  Three times  Four times
4. Are you able to answer the source and background of IMP? Generic  Original
5. Are you able to answer the adverse reaction of the IMP?  
 Dizziness  Headache  Influenza syndrome  Discomfort  Nausea   
 Dyspepsia  Allergy  Angina  Arthralgia
6. Do you know what to do if you feel uncomfortable?  
 Yes  No

**Thank you very much for your filling!**