Ethics Committee of Women’s Hospital, School of Medicine, Zhejiang University

Approval for Ethical Review

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| Research Title | Risk factors of preeclampsia in pregnant women with polycystic ovary syndrome |
| Research source | 口Enterprises, 口government, 口academic organizations,√this unit, 口self-funded, 口others |
| Research Unit | Women’s Hospital, School of Medicine, Zhejiang University |
| Principal Researcher | Jiang Ruoan |
| Study Protocol No. | 1.0 | Date of Protocol Version | 2022-06-30 |
| Informed ConsentNo. | - | Date of Informed Consent Version | - |
| Review Mode | A quick review | Review Site | - |
| Meeting Date | - | Approval NO. | IRB-20220220-R |
| Review Opinion |  |  |  |
| Upon the decision of the ethics committee, we agree to carry out this mantra study according to the approved mantra study plan and other materials.Please follow the principles of GCP and ethics to protect the health and rights of subjects.Note: this ethical approval is only used for scientific research and cannot be used for any commercial activities. The research data cannot be provided to the enterprises that have nothing to do with the research. |
| Tracking review frequency | 12 months | Deadline for approval | 2023-07-14 |
| Signature |  |  |  |
| Ethic committee |  |  |  |

pay attention to:

1 This ethics committee is independent and complies with ICH GCP, Chinese GCP and ethics-related regulations.The committee review and working process are not affected by any organization or individual.

2 This ethics committee will protect the reviewed clinical research data and relevant content. 3 During the study, modify the clinical research and filling case, informed consent and other materials.Please submit the amendment application.

4 When a SAE or a SUSAR occurs, inform the ethics Committee in time.

5 The applicant is requested to submit the research road exhibition report I month before the validity period of the approval according to the annual or regular audit frequency approved by the Ethics Committee.

6 Please promptly report to the ETHICS Committee any situation during the study that may significantly affect the conduct of the study or increase the risk to the subject.

7 In case of any violation of GCP principles during the study, the researcher is requested to submit a violation plan report.

8 Applicants should submit the suspension/termination/completion report in a timely manner when suspending/terminating/completing the study.