**Supplementary Table S2. Detailed inclusion and exclusion criteria for the clinical trial (NCT04370405)**

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| **Inclusion criteria** |
| 1. Voluntarily participating in this clinical trial, understanding the study procedures, and capable of signing an informed consent form in writing;
2. Grade 0-2 ECOG performance status (PS);
3. Expected survival ≥ 3 months;
4. Having at least one measurable disease conforming to IRWG criteria;
5. Good organ function level:
6. ANC ≥ 1.0 × 109/L;
7. PLT ≥ 70 × 109/L (≥ 50 × 109/L for patients with marrow involvement);
8. Hb ≥ 80 g/L (≥ 70 g/L for patients with marrow involvement);
9. TBIL ≤ 1.5 × ULN;
10. ALT and AST ≤ 2.5 × ULN;
11. BUN/Urea and Cr ≤ 1.5 × ULN;
12. LVEF ≥ 50%;
13. Corrected QT interval by Fridericia method (QTcF) < 450 ms for males and < 470 ms for females.
14. Not participated in a clinical trial as a subject within 1 month before the present trial;
15. Capable of following the trial protocol in the investigator’s opinion.
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| **Exclusion criteria** |
| 1. Pleural effusion or ascites that cannot be controlled by drainage or other methods;
2. Various factors affecting drug administration and absorption, such as inability to swallow, chronic diarrhea or intestinal obstruction;
3. For patients who are receiving treatment with medication that may lead to a prolonged QT interval (such as antiarrhythmic drugs), the treatment cannot be discontinued or switched to a different medication prior to starting study drug;
4. Tumor with central nervous system (CNS) invasion;
5. Any cardiac disorders including (1) angina; (2) arrhythmia requiring clinical treatment and intervention; (3) myocardial infarction; (4) cardiac failure; (5) any other cardiac disorders which cause the patients unsuitable to be enrolled in this trial in the investigator’s opinion;
6. Female in gestation period or lactation period, or female with childbearing potential who has positive baseline pregnancy test result;
7. Severe concomitant diseases (e.g. severe hypertension, diabetes, thyroid disorder, etc.) that compromise the safety of patients or prevent patients from completing the study in the investigator’s opinion;
8. Other neoplasm malignancy within the past 5 years, except cured skin basal cell carcinoma and cervix carcinoma in situ;
9. Receipt of an attenuated live vaccine within 30 days before the first dose.
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