

Efficacy and Safety of Prophylactic Treatment for Pneumocystis Jirovecii Pneumonia in Patients With Autoimmune Inflammatory Rheumatic Disease

NCT06499233

Status	RECRUITING
Phase	Phase 4
Sponsor	Tongji Hospital
Enrollment	800 participants

Key Eligibility Criteria

Inclusion (3)

- The patient was diagnosed with AIIRD according to the International Classification of Diseases and had received steroids or immunosuppressive therapy;
- The patient had not received standard PJP treatment before enrollment, including the first-line treatment drug TMP/SMZ, or other second-line treatment drugs (including Pentamidine, Atorvastatin, Caspofungin, etc.);
- The patient was at least 18 years old at the time of enrollment;

Exclusion (7)

- Serious health problems or diseases, including (but not limited to) the following: severe liver damage (ALT, AST elevated above normal value by more than 5 times), severe renal insufficiency (GFR $< 30\text{mL/min}$ or Scr $> 445\mu\text{mol/L}$), severe myelosuppression (Hb $< 65\text{g/L}$, PLT $< 25 \times 10^9/\text{L}$ or neutrophils $< 0.5 \times 10^9/\text{L}$);
 - Screening test indicates infection with human immunodeficiency virus (HIV), history of lymphomatous hyperplasia of the lymphatic tissue or any malignant tumor of any organ system within the past 5 years, or history of organ transplantation;
 - Participants with a history of allergy to sulfonamide drugs, megaloblastic anemia due to folate deficiency;
 - Pregnant and lactating women;
 - Any medical or psychological condition that the investigator believes would interfere with the participant's ability to comply with the protocol or complete the study;
- ... and 2 more (see full listing online)

Locations (1 total)

Tongji Hospital, Wuhan, Hubei, China

<https://clinicaltrials.gov/study/NCT06499233>

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